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A A UNITED STATES SECURITIES AND EXCHANGE COMMISSION WASHINGTON, DC 20549 A FORM 10-Q (Mark One) a~ QUARTERLY REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934 For the quarterly period ended June 30, 2024 OR a~ TRANSITION REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934 For the transition period from to

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Nuvation Bio Inc. (the “Company” or “Nuvation Bio”). On April 9, 2024 (the “Acquisition Date”), the Company completed its acquisition (the “Acquisition”) of AnHeart Therapeutics Ltd., an exempted company incorporated under the laws of the Cayman Islands (the “AnHeart”), pursuant to that certain Agreement and Plan of Merger (the “AnHeart Merger Agreement”), by and among the Company, AnHeart, Artemis Merger Sub I, Ltd., an exempted company incorporated under the laws of the Cayman Islands and a wholly owned subsidiary of the Company, and Artemis Merger Sub II, Ltd., an exempted company incorporated under the laws of the Cayman Islands and a wholly owned subsidiary of the Company. 2. Significant Accounting Policies Basis of Presentation The accompanying consolidated financial statements have been prepared in accordance with accounting principles generally accepted in the United States of America (the “GAAP”) and include all adjustments necessary for the fair presentation of the Company’s financial position for the periods presented. Principles of Consolidation The consolidated financial statements include the balances of the Company and its subsidiaries. All intercompany transactions and balances are eliminated in consolidation. Unaudited Interim Financial Information The accompanying unaudited condensed consolidated financial statements as of June 30, 2024 and for the three and six months ended June 30, 2024 and 2023, have been prepared in accordance with GAAP for interim financial statements and pursuant to Form 10-Q and Article 10 of Regulation S-X. In the opinion of management, all adjustments (consisting of normal recurring accruals) have been made that are considered necessary for a fair statement of the financial position of the Company as of June 30, 2024, the results of operations for the three and six months ended June 30, 2024 and 2023 and the cash flows for the six months ended June 30, 2024 and 2023. The condensed consolidated balance sheet as of December 31, 2023 has been derived from the Company’s audited consolidated financial statements. Operating results for the three and six months ended June 30, 2024 are not necessarily indicative of the results that may be expected for the year ending December 31, 2024. Certain information and disclosures normally included in the notes to annual financial statements prepared in accordance with GAAP have been omitted from these interim financial statements pursuant to the rules and regulations of the Securities and Exchange Commission (the “SEC”). Accordingly, these unaudited condensed consolidated financial statements should be read in conjunction with the financial statements for the fiscal year ended December 31, 2023, which are included in the Company’s 10-K filed with the SEC on February 29, 2024. Liquidity As of June 30, 2024, the Company has an accumulated deficit of approximately \$820.1 million and net cash used in operating activities was approximately \$53.0 million for the six months ended June 30, 2024. Management expects to continue to incur operating losses and negative cash flows from operations for the foreseeable future. A 5 A As of June 30, 2024, the Company had cash, cash equivalents, and marketable securities of \$577.2 million. The Company believes that its existing cash, cash equivalents, and marketable securities will be sufficient to meet its cash commitments for at least the next 12 months after the date that these consolidated financial statements are issued. The Company’s research and development and preparation for potentially becoming a commercial stage organization can be costly, and the timing and outcomes are uncertain. The assumptions upon which the Company has based its estimates are routinely evaluated and may be subject to change. The actual amount of the Company’s expenditures will vary depending upon a number of factors including but not limited to the progress of the Company’s research and development activities and potential regulatory approval of its product candidates and the level of financial resources available. Significant Risks and Uncertainties The Company’s operations are subject to a number of factors that can affect its operating results and financial condition. Such factors include, but are not limited to: the results of research and development, clinical testing and trial activities of the Company’s products, the Company’s ability to obtain regulatory approval to market its products, competition from products manufactured and sold or being developed by other companies, the price of, and demand for, Company’s products, the Company’s ability to negotiate favorable licensing or other manufacturing and marketing agreements for its products, and the Company’s ability to raise capital. The Company currently has no commercially approved products and there can be no assurance that the Company’s research and development will be successfully commercialized. Developing and commercializing a product requires significant time and capital and is subject to regulatory review and approval as well as competition from other biotechnology and pharmaceutical companies. The Company operates in an environment of rapid change and is dependent upon the continued services of its employees and vendors and obtaining and protecting intellectual property. Use of Estimates The preparation of financial statements in conformity with GAAP requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and the disclosure of contingent assets and liabilities at the date of the financial statements and the reported amount of revenue and expenses during the year. Accordingly, actual results could differ from those estimates and those differences could be significant. Significant estimates and assumptions reflected in the accompanying consolidated financial statements include, but are not limited to, warrant liabilities, leases, stock options granted and depreciation expense. Functional Currency and Foreign Currency Translations The Company’s reporting currency is the U.S. dollar (“USD”). The functional currency of the Company’s subsidiaries incorporated in People Republic of China (“PRC”) is Renminbi (“RMB”). The functional currency of the Company and its subsidiaries incorporated outside the PRC is USD. Transactions denominated in currencies other than the functional currencies are re-measured into the functional currency of the entity at the exchange rates prevailing on the transaction dates. Foreign currency denominated financial assets and liabilities are re-measured at the balance sheet date exchange rate. The resulting exchange differences are included in the net loss of the statements of operations and comprehensive loss. Assets and liabilities of the Company with functional currency other than USD are translated into USD at fiscal year-end exchange rates. Equity amounts are translated at historical exchange rates. Income and expense items are translated at average exchange rates during the fiscal year. Translation adjustments arising from these are reported as foreign currency translation adjustments and are shown as a component of other comprehensive income (loss). Cash and Cash Equivalents Cash equivalents include short-term, highly liquid instruments, consisting of money market accounts, a money market mutual fund and short-term investments with maturities from the date of purchase of 90 days or less. The majority of cash and cash equivalents are maintained with major financial institutions in North America. Deposits with these financial institutions may exceed the amount of insurance provided on such deposits. These deposits may be redeemed upon demand which reduces counterparty performance risk. Marketable Securities Debt securities have been classified as available-for-sale which may be sold before maturity or are not classified as held to maturity or trading. Marketable debt securities classified as available-for-sale are carried at fair value with unrealized gains or losses reported in other comprehensive income (loss). Exchange traded funds are equity securities, which are reported as marketable securities with readily determinable fair values, are also carried at fair value with unrealized gains and losses included in other income (expense), net. Realized gains and losses on both debt and equity securities are included in other income (expense), net. For securities in an unrealized loss position, management considers the extent and duration of the unrealized loss, and the financial condition and near-term prospects of the issuer. Management also assesses whether it intends to sell, or it is more likely than not that it will be required to sell, a security in an unrealized loss position before recovery of its amortized cost basis. If management determines there is any other than temporary impairment, the entire difference between amortized cost and fair value is recognized as impairment through earnings. The Company is exposed to credit losses primarily through its available-for-sale investments. The Company assesses whether its available-for-sale investments are impaired at each reporting period. Unrealized losses or impairments resulting from the amortized cost basis of any available-for-sale debt security exceeding its fair value are evaluated for identification of credit losses and non-credit related losses. Any credit losses are charged to earnings against the allowance for credit losses of the debt security, limited to the difference between the fair value and the amortized cost basis of the debt security. Any difference between the fair value of the debt security and the amortized cost basis, less the allowance for expected credit losses, are reported in other comprehensive income (loss). Expected cash inflows due to improvements in credit are recognized through a reversal of the allowance for expected credit losses subject to the total allowance previously recognized. The Company’s expected loss allowance methodology for the debt securities includes reviewing the extent of the unrealized loss, the size, term, geographical location, and industry of the issuer, the issuer’s credit ratings and any changes in those ratings, as well as reviewing current and future economic market conditions and the issuer’s current status and financial condition. As of June 30, 2024, the Company has not recognized an allowance for expected credit losses related to available-for-sale investments as the Company has not identified any unrealized losses for these investments attributable to credit factors. Interest income includes amortization and accretion of purchase premiums and discounts. Premiums and discounts on debt securities are amortized on the effective-interest method. Gains and losses on sales are recorded on the settlement date and determined using the specific identification method. Concentration of Risk Concentration of Risk Financial instruments that potentially subject the Company to concentrations of credit risk are primarily cash, cash equivalents and marketable securities. The Company maintains its cash and cash equivalent balances in the form of business checking accounts and money market accounts, the balances of which, at times, may exceed federally insured limits. Exposure to cash and cash equivalents credit risk is reduced by placing such deposits with major financial institutions and monitoring their credit ratings. Marketable securities consist primarily of government and corporate bonds, municipal securities and exchange traded funds with fixed interest rates. Exposure to credit risk of marketable securities is reduced by maintaining a diverse portfolio and monitoring their credit ratings. Concentration of customers Substantially all revenues were derived from customers located in China and Japan. For the three and six months ended June 30, 2024, the revenues derived from customers located in China and Japan were \$1.1 million and \$0.3 million, respectively. There was no revenue in 2023 since the Acquisition of AnHeart completed in the second quarter of 2024. Below customers represent more than 10% of the Company’s net revenue for the three and six months ended June 30, 2024. A 3 Three Months Ended June 30, A 6 Six Months Ended June 30, A 4 2024 A 4 2024 A Customer A 4 1,096 A 4 1,096 A Customer B A 4 339 A 4 339 A A A Below customer represents more than 10% of the Company’s balances of accounts receivable as of June 30, 2024. A 4 June 30, 2024 A Customer B A 4 117 A A A Property and Equipment Property and equipment are stated at cost less accumulated depreciation. Property and equipment are depreciated using the straight-line method over the estimated useful lives of the related assets of generally five years for computers and seven years for A 7 furniture and equipment. The cost of leasehold improvements is amortized on the straight-line method over the lesser of the estimated asset life or remaining term of the lease. Maintenance costs are expensed as incurred, while major betterments are capitalized. Impairment of Long-Lived Assets The Company reviews long-lived assets for impairment whenever events or changes in circumstances indicate that the carrying amount of an asset may not be recoverable and an impairment assessment may be performed on the recoverability of the carrying amounts. If an impairment occurs, the loss is measured by comparing the fair value of the asset to its carrying amount. Warrants The Company accounts for warrants as either equity-classified or liability-classified instruments based on an assessment of the warrant’s specific terms and applicable authoritative guidance in Financial Accounting Standards Board (the “FASB”) Accounting Standards Codification (the “ASC”) 480, Distinguishing Liabilities from Equity (the “ASC 480”) and ASC 815, Derivatives and Hedging (the “ASC 815”). The assessment considers whether the warrants are freestanding financial instruments pursuant to ASC 480, meet the definition of a liability pursuant to ASC 480, and whether the warrants meet all of the requirements for equity classification under ASC 815, including whether the warrants are indexed to the Company’s own ordinary shares, among other conditions for equity classification. This assessment, which requires the use of professional judgment, is conducted at the time of warrant issuance and as of each subsequent quarterly period end date while the warrants are outstanding. For issued or modified warrants that meet all of the criteria for equity classification, the warrants are required to be recorded as a component of additional paid-in capital at the time of issuance. For issued or modified warrants that do not meet all the criteria for equity classification, the warrants are required to be recorded as a liability at their fair value on the date of issuance and each balance sheet date thereafter. Changes in the estimated fair value of the warrants are recognized as a non-cash gain or loss on the statements of operations and comprehensive loss. Following the Merger, there were 5,787,472 warrants to purchase Common Stock outstanding, consisting of 4,791,639 Public Warrants, 162,500 Private Placement Warrants and 833,333 Forward Purchase Warrants. Each whole warrant entitles the registered holder to purchase one share of our Class A Common Stock at a price of \$11.50 per share. Pursuant to the warrant agreement, a warrant holder may exercise its warrants only for a whole number of shares of our Class A Common Stock. As part of the consideration for AnHeart, the Company issued warrants collectively exercisable for approximately 2,893,731 shares of the Company’s Class A Common Stock at an exercise price of \$11.50 per share (the “Consideration Warrants”). The fair value of the Consideration Warrants was determined to approximate the public warrant trading price as of closing of the Acquisition, yielding a fair value of approximately \$0.26 per share. The Consideration Warrants are restricted with respect to the exercise and transfer thereof until receipt of such stockholder approval and otherwise have terms identical to those of the Company’s outstanding Public Warrants. Accordingly, the Company classified the Consideration Warrants as liabilities, consistent with the classification of the Company’s Public Warrants. The Company evaluated Public Warrants, Private Placement Warrants, Consideration Warrants, and Forward Purchase Warrants (the “Warrants”) under ASC 815-40, Derivatives and Hedging’s “Contracts in Entity’s Own Equity,” and concluded that they do not meet the criteria to be classified in stockholders’ equity. Specifically, the settlement value of the Warrants is dependent, in part, on the holder of the Warrants at the time of settlement. Because the holder of an instrument is not an input into the pricing of a fixed-for-fixed option on our Common Stock, the Warrants fail the indexation guidance in ASC 815-40, which would preclude classification in stockholders’ equity. Additionally, the exercise of the Warrants may be settled in cash upon the occurrence of a tender offer or exchange that involves more than 50% of the outstanding shares of the Company’s Common Stock. Because not all of the Company’s stockholders need to participate in such tender offer or exchange to trigger the potential cash settlement and the Company does not control the occurrence of such an event, the Company concluded that the Warrants do not meet the conditions to be classified in equity. Since the Warrants meet the definition of a derivative under ASC 815, the Company recorded these Warrants as liabilities on the balance sheet at fair value upon the closing of the Merger, with subsequent changes in their respective fair values recognized in the consolidated statement of operations and comprehensive loss at each reporting date. The fair value of Consideration, Public, and Forward Purchase Warrants was determined using the closing price of the publicly traded warrants on the NYSE market. The fair value of the Private Placement Warrants was estimated using a Black-Scholes option pricing model (see Note 4). Revenue Recognition The Company applies ASC Topic 606, “Revenue from Contracts with Customers” (“ASC 606”) to account for its revenue transactions. Under ASC 606, an entity recognizes revenue when its customer obtains control of promised goods or services, in an amount that reflects the consideration that the entity expects to receive in exchange for those goods or services. To determine revenue recognition for arrangements that an entity determines are within the scope of ASC 606, the entity performs the five-step model. The Company only applies the five-step model to contracts when it is probable that the entity will collect the consideration to which it is entitled in exchange for the goods or services it transfers to the customer. Once a contract is determined to be within the scope of ASC 8 A 606 at contract inception, the Company reviews the contract to determine which performance obligations it must deliver and which of these performance obligations are distinct. The Company recognizes as revenue the amount of the transaction price that is allocated to each performance obligation when that performance obligation is satisfied or as it is satisfied. Accounts receivable represent amounts invoiced and revenues recognized prior to invoicing when the Company has satisfied its performance obligation and has the unconditional right to payment. Collaborative Arrangement In November 2018, the FASB issued Topic 808, “Collaborative Arrangements” (“ASC 808”): Clarifying the Interaction between ASC 808 and ASC 606. This update clarifies that certain transactions between participants in a collaborative arrangement should be accounted for under ASC 606 when the counterparty is a customer and precludes an entity from presenting consideration from a transaction in a collaborative arrangement as revenue from contracts with customers if the counterparty is not a customer for that transaction. The Company adopted this standard for all periods presented. The Company enters into collaborative arrangements for the research and development, manufacture and/or commercialization of drug products and drug candidates. The Company assessed and determined that none of the collaboration agreements entered into during the periods presented were within the scope of ASC 808, as all of the agreements did not involve active participation by both parties in a joint research activity, and therefore did not qualify as collaborative arrangements under ASC 808. The Company has determined that all the elements of the above collaborations are reflective of a vendor-customer relationship and therefore within the scope of ASC 606. In determining the appropriate amount of revenue to be recognized as it fulfills its obligations under each of its agreements, the Company performs the five-step model under ASC 606 noted above. The Company recognizes revenue to depict the transfer of control of promised goods or services to customers in an amount that reflects the consideration to which the entity expects to receive in exchange for those goods or services. The Company’s collaborative arrangements may contain more than one unit of account, or performance obligation, including grants of licenses to intellectual property rights, agreements to provide research and development services and other deliverables. The collaborative arrangements do not include a right of return for any deliverable. As part of the accounting for these arrangements, the Company must develop assumptions that require judgment to determine the stand-alone selling price for each performance obligation identified in the contract. In developing the stand-alone selling price for a performance obligation, the Company considers competitor pricing for a similar or identical product, market awareness of and perception of the product, expected product life and current market trends. In general, the consideration allocated to each performance obligation is recognized when the respective obligation is satisfied either by delivering a good or providing a service, limited to the consideration that is not constrained. Non-refundable payments received before all of the relevant criteria for revenue recognition are satisfied are recorded as contract liabilities. Licenses of intellectual

provides a license to the Company's intellectual property which is determined to be distinct from the other promises or performance obligations identified in the contract, the Company recognizes revenue from the portion of the transaction price allocated to the license at a point in time, when the license is transferred to the customer and the customer is able to benefit from the license. Research and development servicesThe portion of a transaction price allocated to research and development services performance obligations is deferred and recognized as revenue over time as delivery or performance of such services occurs based on the use of an input method. Customer optionsA customer's right to choose, at its discretion, to make a payment for additional goods or services is generally considered an option. If the Company is not presently obligated to provide and does not have a right to consideration for delivering additional goods or services, the item is considered an option. The Company evaluates the customer options for material rights, such as the ability to acquire additional goods or services for free or a discount. Optional future services that reflect their standalone selling prices do not provide the customer with a material right and, therefore, are not considered performance obligations and are accounted for as separate contracts. The optional future services do not include a material right to be accounted for as performance obligations. Milestones paymentsThe Company's collaboration agreements include development and regulatory milestones. The Company evaluates whether the milestones are considered probable of being reached and estimates the amount to be included in the transaction price using the most likely amount method. The Company evaluates factors such as the scientific, clinical, regulatory, commercial, and other risks that must be overcome to achieve the particular milestone in making this assessment. If it is probable that a significant revenue reversal would not occur, the associated milestone value is included in the transaction price. Milestone payments that are not within the Company's control or the licensee's control, such as regulatory approvals, are not considered probable of being achieved until those approvals are received. At the end of each reporting period, the Company re-evaluates the probability of achievement of such milestones and any related ASC 940-608 constraint, and if necessary, adjusts the estimate of the overall transaction price. Any such adjustments are recorded on a cumulative catch-up basis, which would affect revenue and net loss in the period of adjustment. RoyaltiesFor sales-based royalties, including milestone payments based on the level of sales, the Company determines whether the sole or predominant item to which the royalties relate is a license. When the license is the sole or predominant item to which the sales-based royalty relates, the Company recognizes revenue at the later of: (i) when the related sales occur, or (ii) when the performance obligation to which some or all of the royalty has been allocated has been satisfied (or partially satisfied). To date, the Company has not recognized any sales-based royalty revenue resulting from the Company's collaboration agreements. The Company receives payments from its customers based on billing terms established in the contract. Up-front payments and fees are recorded as contract liabilities (e.g., deferred revenue) upon receipt or when due until the Company performs its obligations under the arrangement. In the event of an early termination of a collaboration agreement, any contract liabilities would be recognized in the period in which all our obligations under the agreement have been fulfilled. For a complete discussion of accounting for revenue, see Note 5, "Collaboration and License Agreements."Costs to Fulfill a Contract with a CustomerThe Company has not capitalized any costs for the three and six months ended June 30, 2024 and 2023. The Company is required to capitalize costs incurred to fulfill customer contracts. These costs are required to be amortized to expense on a systemic basis that is consistent with the transfer to the customer of the goods or services to which the asset relates, compared to previously being expensed as incurred. Elements of the costs primarily include (i) payroll and other related costs of personnel related directly to the contract activities; (ii) costs related to pre-clinical testing and clinical trials such as payments to contract research organizations ("CROs") and contract manufacturing organizations ("CMOs"), investigators, and clinical trial sites that conduct the contract activities; (iv) costs to develop the product candidates, including raw materials and supplies, product testing, depreciation, and facility-related expenses; and (v) other research and development costs.LeasesThe Company determines if an arrangement contains a lease at inception. For arrangements where the Company is the lessee, operating leases are included in operating lease right-of-use, or ROU assets; current operating lease liabilities; and non-current operating lease liabilities on its balance sheets. The Company currently does not have any finance leases.Operating lease ROU assets and operating lease liabilities are recognized based on the present value of the future minimum lease payments over the lease term at the commencement date. ROU assets also include any initial direct costs incurred and any lease payments made on or before the lease commencement date, less lease incentives received. The Company uses its incremental borrowing rate based on the information available at the commencement date in determining the lease liabilities as the Company's leases generally do not provide an implicit rate. The incremental borrowing rate is reevaluated upon a lease modification. The operating lease ROU asset also includes any initial direct costs and prepaid lease payments made less any lease incentives. The Company considered information available at the adoption date of Topic 842 to determine the incremental borrowing rate for leases in existence as of this date. The incremental borrowing rate used was the weighted average rate between secured and unsecured lending arrangements. Lease terms may include options to extend or terminate the lease when the Company is reasonably certain that the option will be exercised. Variable payments included in the lease agreement are expensed as incurred. Lease expense is recognized on a straight-line basis over the lease term.The Company elected to apply each of the practical expedients described in ASC Topic 842-10-65-1(f) which allow companies not to reassess: (i) whether any expired or existing agreements contain leases, (ii) the classification of any expired or existing leases, and (iii) the capitalization of initial direct costs for any existing leases. The Company also elected to apply the short-term lease measurement and recognition exemption in which ROU assets and lease liabilities are not recognized for short-term operating leases. A short-term operating lease is a lease that, at the commencement date, has a lease term of 12 months or less and does not include an option to purchase the underlying asset that the lessee is reasonably certain to exercise. The Company also elected not to separate non-lease components from lease components and instead to account for each separate lease component and the non-lease components associated with that lease component as a single lease component.Segment InformationThe Company manages its operations as a single segment for the purposes of assessing performance and making operating decisions. The Company's operations are focused on oncology development activities.A 10 Research and Development CostsCosts incurred in connection with research and development activities are expensed as incurred. These costs include fees paid to consultants, vendors and various entities that perform certain research and testing on behalf of the Company. The Company has acquired rights to develop and commercialize product candidates. Prepaid payments that relate to the acquisition of a new product compound, as well as pre-commercial milestone payments, are immediately expensed as acquired in-process research and development in the period in which they are incurred, provided that the new product compound did not also include processes or activities that would constitute a "business" as defined under U.S. GAAP, and the product candidate has not achieved regulatory approval for marketing and, absent obtaining such approval, has no established alternative future use. Milestone payments made to third parties subsequent to regulatory approval which meet the capitalization criteria would be capitalized as intangible assets and amortized over the estimated remaining useful life of the related product. If the conditions enabling capitalization of development costs as an asset have not yet been met, all development expenditures are recognized in profit or loss when incurred.Stock-based CompensationThe Company recognizes compensation cost for grants of employee stock options using a fair-value measurement method, that is recognized in operating results as compensation expense based on the fair value over the requisite service period of the awards. Forfeitures are recorded as they occur instead of estimating forfeitures that are expected to occur.The Company determines the fair value of stock-based awards that are based only on a service condition using the Black-Scholes option-pricing model which uses both historical and current market data to estimate fair value. The method incorporates various assumptions such as the risk-free interest rate, volatility, dividend yield, and expected life of the options. The Company determines the fair value of stock-based awards that are based on both a service condition and achievement of the first to occur of a market or performance condition using a Monte Carlo simulation.Income TaxesThe Company accounts for income taxes using the asset and liability method, which requires the recognition of deferred tax assets and liabilities for the expected future tax consequences of events that have been recognized in the consolidated financial statements or in the Company's tax returns. The difference between the financial statement and tax basis of assets and liabilities is determined annually. Deferred income tax assets and liabilities are computed for those differences that have future tax consequences using the currently enacted tax laws and rates that apply to the years in which they are expected to affect taxable income. The Company assesses the likelihood that its deferred tax assets will be recovered from future taxable income and, to the extent it believes, based upon the weight of available evidence, that it is more likely than not that all or a portion of the deferred tax assets will not be realized, a valuation allowance is established through a charge to income tax expense.The Company uses a recognition threshold and measurement attribute for the financial statement recognition and measurement of a tax position taken, or expected to be taken, in a tax return. The provision for income taxes includes the effects of any resulting tax reserves, or unrecognized tax benefits, that are considered appropriate. The Company's policy is to record interest and penalties related to income taxes as part of the tax provision. Returns for tax years beginning with those filed for the period ended December 31, 2018 are open to federal and state tax examination.Recent Accounting PronouncementsIn November 2023, the FASB issued Topic 280 "Improvements to Reportable Segment Disclosures" which expands annual and interim disclosure requirements for reportable segments, primarily through enhanced disclosures about significant segment expenses. This guidance is effective for our annual periods beginning January 1, 2024, and for interim periods beginning January 1, 2025, with early adoption permitted. We are currently evaluating the potential effect that the updated standard will have on our financial statement disclosures.In December 2023, the FASB issued Topics 740 "Improvements to Income Tax Disclosures" to expand the disclosure requirements for income taxes, specifically related to the rate reconciliation and income taxes paid. This guidance is effective for our annual periods beginning January 1, 2025, with early adoption permitted. We are currently evaluating the potential effect that the updated standard will have on our financial statement disclosures. 3. AcquisitionOn the Acquisition Date, the Company completed the Acquisition of AnHeart. Pursuant to the terms of the AnHeart Merger Agreement, at the effective time of the First Merger (as defined in the Merger Agreement) (the "First Effective Time"), the Company issued to AnHeart securityholders (i) approximately 27,646,255 shares of Class A Common Stock, (ii) 851,202 shares of Series A Non-Voting Convertible Preferred Stock of the Company, and (iii) the Consideration Warrants. The shares of our Series A Non-Voting Convertible Preferred Stock are automatically convertible into an aggregate of approximately 85,120,200 shares of Class A Common Stock upon the approval of such conversion by the Company's stockholders in accordance with the rules of the New York Stock Exchange. Since the preferred stock is not mandatorily redeemable and the exception for a deemed liquidation event was not met, all of the outstanding convertible preferred stock is classified as mezzanine equity in the balance sheet. The Consideration Warrants are restricted with respect to the exercise and transfer thereof until receipt of such stockholder approval and otherwise have terms identical to those of the Company's outstanding Public Warrants.At the First Effective Time, (i) each option to purchase shares of Ordinary Shares (as defined in the Merger Agreement) of AnHeart (an "AnHeart Option") held by a Continuing Company Service Provider (as defined in the Merger Agreement), whether or not vested, was assumed and converted into an option to purchase Class A Common Stock (each such option, an "Assumed Option"), and (ii) each restricted stock unit reflecting the right to receive Ordinary Shares of AnHeart (an "AnHeart RSU") held by a Continuing Company Service Provider (as defined in the Merger Agreement) was assumed and became a restricted stock unit with respect to a number of shares of Class A Common Stock (each such restricted stock unit, an "Assumed RSU"). Each Assumed Option and Assumed RSU is subject to the same terms and conditions (including vesting and exercise schedule) as were applicable to the corresponding AnHeart Option or AnHeart RSU immediately prior to the First Effective Time, subject to limited exceptions set forth in the Merger Agreement. Any other AnHeart Option and AnHeart RSU that remained unexercised and outstanding as of immediately prior to the First Effective Time was canceled without payment. From and after the Effective Time, the number of shares of Class A Common Stock subject to each Assumed Option was determined by multiplying (A) the number of Ordinary Shares of AnHeart that are subject to such AnHeart Option immediately prior to the First Effective Time, by (B) the Equity Award Exchange Ratio (as defined in the Merger Agreement) and rounding down to the nearest whole number of shares; the per-share exercise price for Class A Common Stock issuable upon exercise of each Assumed Option was determined by dividing (A) the per-share exercise price of each AnHeart Option, as in effect immediately prior to the First Effective Time, by (B) the Equity Award Exchange Ratio and rounding up to the nearest whole cent; and the number of shares of Class A Common Stock subject to each Assumed RSU was determined by multiplying (A) the number of Ordinary Shares of AnHeart that are subject to such AnHeart RSU immediately prior to the First Effective Time, by (B) 1.7551 and rounding down to the nearest whole number of shares. The final allocable purchase price consists of the following (in thousands): Class A Common Stock (1) \$ 89,297.1 Series A Non-Voting Convertible Preferred Stock (2) \$ 274,938.4 Consideration Warrants (3) \$ 764.4 Assumed share-based awards (4) \$ 24,818.4 Acquisition costs \$ 7,434.4 Total allocable purchase price \$ 397,251.4 \$ 27,646,255 shares issued at \$3.23 per share as of the Acquisition Date. (2) 851,202 shares issued at as-converted value of \$323 per share.(3) 2,893,731 warrants issued at the estimated fair value of \$0.2641 per warrant.(4) The portion recorded in consideration transferred represents the fair value of replacement awards (Assumed Options & Assumed RSUs) attributable to precombination services.We accounted for the transaction as an asset acquisition since the lead asset, tislelizumab, represented substantially all of the fair value of the gross assets acquired, which exclude cash acquired. At the Acquisition Date, we recorded a \$425.1 million charge representing an acquired in-process research and development asset with no alternative future use in acquired in-process research and development expenses, of which the \$425.1 million net cash consideration is presented as a cash outflow from operating activities. In connection with this acquisition, we recorded \$450.9 million of assets acquired primarily consisting of cash and investments. Liabilities assumed were approximately \$53.7 million.

4. Fair Value Measurements and Marketable Securities Available-for-SaleThe Company provides disclosure of financial assets and financial liabilities that are carried at fair value based on the price that would be received upon sale of an asset or paid to transfer a liability in an orderly transaction between market participants at the measurement date. Fair value measurements may be classified based on the amount of subjectivity associated with the inputs to fair valuation of these assets and liabilities using the following three levels:Level 1 Inputs are unadjusted quoted prices in active markets for identical assets or liabilities that the Company has the ability to access at the measurement date.Level 2 Inputs include quoted prices for similar assets and liabilities in active markets, quoted prices for identical or similar assets or liabilities in markets that are not active, inputs other than quoted prices that are observable for the asset or liability (i.e., interest rates, yield curves, etc.) and inputs that are derived principally from or corroborated by observable market data by correlation or other means (market corroborated inputs).Level 3 Unobservable inputs that reflect the Company's estimates of the assumptions that market participants would use in pricing the asset or liability. The Company develops these inputs based on the best information available, including its own data.The following table presents information about the Company's marketable securities as of June 30, 2024 and December 31, 2023 and the Warrant liability as of June 30, 2024 and December 31, 2023, measured at fair value on a recurring basis, and indicates the fair value hierarchy of the valuation techniques the Company utilized to determine such fair value. The Company's Warrant liabilities are included within the Level 1 and Level 3 fair value hierarchy. The fair value of the Consideration, Public and Forward Purchase Warrants is determined using the closing price of the warrants on the NYSE market. The fair value of the Private Placement Warrants is determined using the Black-Scholes option pricing formula. The primary unobservable input utilized in determining the fair value of the Private Warrants is the expected volatility. The expected volatility was estimated considering observable Public Warrant pricing, the Company's own historical volatility and the volatility of guideline public companies. There have not been any transfers between the levels during the periods.

	June 30, 2024	December 31, 2023
Cash equivalents:	\$ 4,988.4	\$ 4,988.4
Corporate bonds	\$ 21,694.4	\$ 21,694.4
Certificate of deposits	\$ 408,996.4	\$ 408,996.4
Total financial assets:	\$ 562,440.4	\$ 749,242.4
Financial liabilities:	\$ 1,363.4	\$ 78.4
Cash equivalents:		

[illegible]

increase on January 1st of each year, beginning on January 1, 2022 and continuing through and including January 1, 2031, by 1.0% of the total number of shares of Class A Common Stock and Class B Common Stock outstanding Æ 19 Æ or issuable upon conversion or exercise of outstanding instruments on December 31st of the preceding calendar year or such lesser number of shares of Class A Common Stock as determined by our board of directors. Shares subject to purchase rights granted under the 2021 ESPP that terminate without having been exercised in full will not reduce the number of shares available for issuance under the 2021 ESPP. The stock-based compensation expense included in the Company's condensed statement of operations and comprehensive loss for the three and six months ended June 30, 2024 and 2023 is as follows (in thousands): Æ Æ Three Months Ended June 30, Æ Æ Six Months Ended June 30, Æ Æ 2024 Æ Æ 2023 Æ Æ 2024 Æ Æ 2023 Æ Research and development Æ Æ 4,190 Æ Æ 2,420 Æ Æ 6,482 Æ Æ 4,431 Æ General and administrative Æ Æ 5,379 Æ Æ 3,103 Æ Æ 8,128 Æ Æ 5,929 Æ Æ 9,569 Æ Æ 5,523 Æ Æ 14,610 Æ Æ 10,360 Æ Æ The following table summarizes stock option activity under the 2019 Plan, 2021 Plan and assumed options in the AnHeart Acquisition for the six months ended June 30, 2024 Æ Æ Æ Shares Æ Outstanding at December 31, 2023 Æ Æ 30,649,239 Æ Options assumed for the Acquisition of AnHeart Æ Æ 13,742,239 Æ Granted Æ Æ 11,322,679 Æ Exercised Æ Æ (1,798,190) Æ Forfeited or canceled Æ Æ (1,223,019) Æ Outstanding at June 30, 2024 Æ Æ 52,692,948 Æ Exercisable at June 30, 2024 Æ Æ 23,909,107 Æ Æ The weighted average exercise price of all outstanding options as of June 30, 2024 was \$2.80 per share. The following table summarizes the activity for the restricted stock units assumed in the AnHeart Acquisition for the six months ended June 30, 2024. Æ Æ Restricted Stock Æ Æ Units Æ Non-vested at December 31, 2023 Æ Æ æ" Æ RSUs assumed for the Acquisition of AnHeart Æ Æ 2,201,694 Æ Granted Æ Æ æ" Æ Vested Æ Æ (40,342) Æ Forfeited or canceled Æ Æ æ" Æ Non-vested at June 30, 2024 Æ Æ 2,161,352 Æ Æ Æ Æ 11. Warrants Æ Following the Merger, there were 5,787,472 warrants to purchase Common Stock outstanding, consisting of 4,791,639 Public Warrants, 162,500 Private Placement Warrants and 833,333 Forward Purchase Warrants. Each whole warrant entitles the registered holder to purchase one share of our Class A Common Stock at a price of \$11.50 per share. Pursuant to the warrant agreement, a warrant holder may exercise its warrants only for a whole number of shares of our Class A Common Stock. Following the Acquisition, the Company issued 2,893,731 Consideration Warrants. The Consideration Warrants are restricted with respect to the exercise and transfer thereof until receipt of such stockholder approval and otherwise have terms identical to those of the Company's outstanding Public Warrants. At June 30, 2024, there were an aggregate of 8,681,193 warrants outstanding. The Company concluded that the Consideration Warrants, Public Warrants, Private Warrants and Forward Purchase Warrants do not meet the conditions to be classified in equity. The warrants were recorded at fair value with subsequent changes in fair value reflected in earnings (see Note 4). The change in fair value resulted in a gain of \$1.1 million during the three months ended June 30, 2024. Æ 20 Æ The fair value of Consideration, Public and Forward Purchase Warrants is determined using the closing price of the warrants on the NYSE market and the related Warrant liability is included in Level 1 fair value measurements. The Company utilizes the Black-Scholes option pricing formula to determine the fair value of the Private Warrants at each reporting period, with changes in fair value recognized in the statement of operations. The estimated fair value of the warrant liability for the Private Warrants is determined using Level 3 inputs. Inherent in a binomial options pricing model are assumptions related to expected share-price volatility, expected life, risk-free interest rate and dividend yield. The annualized volatility of the Warrant was based on a calibration to the publicly traded warrant price as of the valuation date. The risk-free interest rate was estimated using linear interpolation assuming a term consistent with the time until the warrants expire, and yield information was based on U.S. Treasury Constant Maturities. The expected life of the warrants is assumed to be equivalent to their remaining contractual term. The dividend rate is based on the historical rate, which the Company anticipates remaining at zero. The aforementioned warrant liabilities are not subject to qualified hedge accounting. There were no transfers between Levels 1, 2 or 3 during the period ended June 30, 2024. The following table provides quantitative assumptions regarding Level 3 fair value measurements: Æ Æ June 30, 2024 Æ Æ December 31, 2023 Æ Stock price Æ Æ \$ 2.92 Æ Æ \$ 1.51 Æ Strike price Æ Æ \$ 11.50 Æ Æ \$ 11.50 Æ Term (in years) Æ Æ 1.6 Æ Æ 2.1 Æ Volatility Æ Æ 98.1 % Æ Æ 82.7 % Risk-free rate Æ Æ 4.9 % Æ Æ 4.1 % Dividend yield Æ Æ 0.0 % Æ Æ 0.0 % Æ The Company determined the following fair values for the outstanding warrants (in thousands): Æ Æ June 30, 2024 Æ Public Warrants Æ Æ 767 Æ Private Placement Warrants Æ Æ 78 Æ Forward Purchase Warrants Æ Æ 133 Æ Consideration Warrants Æ Æ 463 Æ Total Æ Æ 1,441 Æ Æ The following presents changes in liabilities classified in Level 3 of the fair value hierarchy for the six months ended June 30, 2024 (in thousands): Æ Six Months Ended Æ June 30, 2024 Æ Beginning balance Æ Æ \$ 15 Æ Change in fair value of Private Warrants liability recognized in earnings Æ Æ 63 Æ Ending balance Æ Æ 78 Æ Æ Æ Æ 12. Commitments and Contingencies Æ Commitments Æ The Company leases its office space under non-cancellable operating lease agreements. These leases also require the Company to pay real estate taxes and other operational expenses associated with the leased locations and are included in rent expense. The effect of graduating rents, net of the rent credits, is being amortized over the life of the leases so as to result in equal monthly rent expense over the lease term. The Company has standby letters of credit with banks in the aggregate amount of \$0.6 million which serve as security for the New York and San Francisco spaces operating leases. The standby letters of credit automatically renew annually. Æ Contingencies Æ 21 Æ From time to time, the Company is involved in routine litigation that arises in the ordinary course of business. There are no pending significant legal proceedings to which the Company is a party, for which management believes the ultimate outcome would have a material adverse effect on the Company's financial position. Æ Item 2. Management's Discussion and Analysis of Financial Condition and Results of Operations. The following discussion and analysis provide information which our management believes is relevant to an assessment and understanding of our consolidated results of operations and financial condition. You should read the following discussion and analysis of our financial condition and results of operations together with our consolidated financial statements and notes thereto included elsewhere in this report. In addition to historical financial information, this discussion contains forward-looking statements based upon our current expectations that involve risks and uncertainties. Our actual results could differ materially from those anticipated in these forward-looking statements as a result of various factors, including those set forth in the section titled "Item 1A. Risk Factors." Æ Overview Æ We are a late clinical-stage, global biopharmaceutical company tackling some of the greatest unmet needs in oncology by developing differentiated and novel product candidates. We were founded in 2018 by our chief executive officer, David Hung, M.D., who founded Medivation, Inc. and led its successful development of oncology drugs Xtandi® and talazoparib (now marketed as Talzenna®), leading to its \$14.3 billion sale to Pfizer Inc. (ÆPfizerÆ) in 2016. We leverage our team's extensive expertise in medicinal chemistry, preclinical development, drug development, business development, manufacturing, and commercialization to pursue oncology targets validated by strong clinical or preclinical data and develop novel small molecules that improve the activity and overcome the liabilities of currently marketed drugs. As a result of our April 2024 Acquisition of AnHeart Therapeutics Ltd. ("AnHeart"), our most advanced clinical-stage product candidate, taletrectinib, is an oral, potent, central nervous system-active, selective, next-generation ROS1 inhibitor specifically designed for the treatment of patients with ROS1-positive non-small cell lung cancer ("NSCLC"). Taletrectinib is being evaluated for the treatment of patients with advanced ROS1-positive NSCLC in two Phase 2 single-arm pivotal studies: TRUST-I in China, and TRUST-II, a global study. Taletrectinib has been granted Orphan Drug Designation by the U.S. Food and Drug Administration (ÆFDAÆ) for the treatment of patients with ROS1-positive NSCLC and other NSCLC indications, and Breakthrough Therapy Designations by both the U.S. FDA and China's National Medical Products Administration ("NMPA") for the treatment of patients with advanced or metastatic ROS1-positive NSCLC. Based on results of the TRUST-I clinical study, China's NMPA has accepted and granted Priority Review Designations to New Drug Applications for taletrectinib for the treatment of adult patients with locally advanced or metastatic ROS1-positive NSCLC who either have or have not previously been treated with ROS1 tyrosine kinase inhibitors ("TKIs"). Worldwide development and commercial rights to taletrectinib have been in-licensed from Daiichi Sankyo. Commercial rights to taletrectinib have been out-licensed in China and Japan. In addition to taletrectinib, our clinical-stage pipeline includes differentiated, novel oncology product candidates that have been generated from our proprietary drug discovery and development programs or acquired through business development activities: Æ Safusidenib is a novel, oral, potent, brain penetrant, targeted inhibitor of mutant isocitrate dehydrogenase 1 ("IDH1"). Safusidenib is being evaluated in a global Phase 2 study for the treatment of patients with diffuse IDH1-mutant glioma. Æ NUV-1511 is our first clinical-stage drug-drug conjugate ("DDC"), which uses a targeting agent to a widely used chemotherapy agent. NUV-1511 is being evaluated in a Phase 1/2 study for the treatment of patients with advanced solid tumors who previously received and progressed on or after treatment with Enhertu® and/or Tredvela® per approved U.S. FDA labeling, human epidermal growth factor receptor 2-negative ("HER2-") metastatic breast cancer, metastatic castration-resistant prostate cancer (ÆmCRPCÆ), advanced pancreatic cancer, and platinum-resistant ovarian cancer ("PROC"). Æ NUV-868 is a BD2-selective, oral, small molecule bromodomain and extra-terminal ("BET") inhibitor that inhibits BRD4. NUV-868 has completed a Phase 1 monotherapy study for the treatment of patients with advanced solid tumors and a Phase 1b study in combination with olaparib for the treatment of patients with ovarian cancer, pancreatic cancer, mCRPC, triple negative breast cancer, and other solid tumors, and in combination with enzalutamide for the treatment of patients with mCRPC. After an internal analysis of efficacy and safety data collected from these studies, we have decided not to initiate a Phase 2 study of NUV-868 as a monotherapy or in combination with olaparib or enzalutamide in the advanced solid tumor indications that were part of the Phase 1 and Phase 1b study designs. We are evaluating next steps for the NUV-868 program, including further development in combination with approved products for indications in which BD2-selective BET inhibitors may improve outcomes for patients. Recent Developments Æ On April 9, 2024, we completed the previously announced Acquisition of AnHeart. Æ 22 Æ Æ In June 2024, we announced data from our pivotal Phase 2 TRUST-I study evaluating taletrectinib in patients in China, which was published in the Journal of Clinical Oncology (Zhou et al., 2024) and presented at the American Society of Clinical Oncology Annual Meeting (Li et al., 2024). Æ In July 2024, we announced that we will present pooled data from the pivotal Phase 2 TRUST-I and TRUST-II studies of taletrectinib for the treatment of patients with advanced ROS1-positive NSCLC at the European Society of Medical Oncology Congress 2024, which will support our planned New Drug Application in the United States. Æ In July 2024, we announced that taletrectinib has been granted Orphan Drug Designation by the U.S. FDA for the treatment of ROS1-positive NSCLC and other NSCLC indications. Æ In July 2024, we terminated our exclusive license agreement with NewG Lab Co., Ltd. (ÆNewG LabÆ). AnHeart previously entered into an agreement with NewG Lab to develop and commercialize taletrectinib in Korea. Æ We have completed an internal analysis of efficacy and safety data collected from the Phase 1 monotherapy and Phase 1b combination studies of NUV-868. Following this analysis, we have decided not to initiate a Phase 2 study of NUV-868 as a monotherapy or in combination with olaparib or enzalutamide in the advanced solid tumor indications that were part of the Phase 1 and Phase 1b study designs. We are evaluating next steps for the NUV-868 program, including further development in combination with approved products for indications in which BD2-selective BET inhibitors may improve outcomes for patients. Financial Overview Since our inception in 2018, we have focused substantially all of our resources on conducting research and development activities, including drug discovery, preclinical studies, clinical trials, establishing and maintaining our intellectual property portfolio, developing our manufacturing network and managing the manufacture of clinical and research material, hiring personnel, raising capital and providing general and administrative support for these operations. The Company's revenue related to its out-licensing collaborative agreements consist of upfront license fees and research and development services revenue from its collaboration agreements. We have funded our operations to date primarily from the issuance and sale of our common and preferred stock, including through the Merger and a Private Investment in Public Equity ("PIPE") financing in connection with the Merger. We have incurred net losses in each year since inception. As of June 30, 2024, we had an accumulated deficit of \$820.1 million. Substantially all of our net losses have resulted from costs incurred in connection with our research and development programs and from general and administrative costs associated with our operations. We expect to continue to incur significant expenses and increasing operating losses over at least the next several years. We expect our expenses will increase substantially in connection with our ongoing activities, as we: Æ advance product candidates through clinical trials; Æ pursue regulatory approval of product candidates; Æ operate as a public company; Æ continue our preclinical programs and clinical development efforts; Æ continue research activities for the discovery of new product candidates; and Æ manufacture supplies for our preclinical studies and clinical trials. In addition, we expect to incur additional costs associated with operating as a public company, including significant legal, audit, accounting, regulatory, tax-related, director and officer insurance, investor relations and other expenses that we did not incur as a private company. As a result, we will need substantial additional funding to support our continuing operations and pursue our growth strategy. Until such time as we can generate significant revenue from product sales, if ever, we expect to finance our operations through the public or private sale of equity, government or private party grants, debt financings or other capital sources, including potential collaborations with other companies or other strategic transactions. If we are unable to obtain additional funding, we could be forced to delay, reduce or eliminate some or all of our research and development programs, product portfolio expansion or any commercialization efforts, which could adversely affect our business prospects, or we may be unable to continue operations. If we raise funds through strategic collaborations or other similar arrangements with third parties, we may have to relinquish valuable rights to our platform technology, future revenue streams, research programs or product candidates or may have to grant licenses on terms that may not be favorable to us and/or may reduce the value of our Common Stock. Our ability to raise additional funds may be adversely impacted by potential worsening global economic conditions and disruptions to and volatility in the credit and financial markets in the United States and worldwide. Because of the numerous risks and uncertainties associated with product development, we cannot predict the timing or amount of increased expenses or when or if we will be able to achieve or maintain profitability. Æ 23 Æ Components of Results of Operations Revenues The Company enters into collaborative arrangements for the research and development, and commercialization of drug products and drug candidates. To date, these collaborative arrangements have included out-licenses of and options to out-license in-licensed compound to other parties and in-licenses of compound from other parties. These arrangements may include non-refundable upfront payments, contingent obligations for potential development, regulatory and commercial performance milestone payments, cost reimbursement arrangements and royalty payments. The Company receives payments from its customers based on billing terms established in the contract. Up-front payments and fees are recorded as contract liabilities (e.g., deferred revenue) upon receipt or when due until the Company performs its obligations under the arrangement. In the event of an early termination of a collaboration agreement, any contract liabilities would be recognized in the period in which all our obligations under the agreement have been fulfilled. Research and Development Expenses Research and development expenses include: Æ expenses incurred under agreements with third-party contract organizations, and consultants; Æ costs related to production of drug substance, including fees paid to contract manufacturers; Æ laboratory and vendor expenses related to the execution of preclinical trials; and Æ employee-related expenses, which include salaries, benefits and stock-based compensation. We expense all research and development costs in the periods in which they are incurred. Costs for certain development activities are recognized based on an evaluation of the progress to completion of specific tasks and estimates of services performed using information and data provided to us by our vendors and third-party service providers. Nonrefundable advance payments for goods or services to be received in future periods for use in research and development activities are deferred and capitalized. The capitalized amounts are then expensed as the related goods are delivered and as services are performed. We expense in-process research and development projects acquired as part of asset acquisitions that have no alternative future use. We expect our research and development expenses to increase substantially for the foreseeable future as we continue to invest in research and development activities related to developing our product candidates, as our product candidates advance into later stages of development, and as we begin to conduct clinical trials. The process of conducting the necessary clinical research to obtain regulatory approval is costly and time-consuming, and the successful development of our product candidates is highly uncertain. As a result, we are unable to determine the duration and completion costs of our research and development projects or when and to what extent we will generate revenue from the commercialization and sale of any of our product candidates. General and Administrative Expenses General and administrative expenses consist of personnel-related costs, facilities costs, depreciation and amortization expenses and professional services expenses, including legal, human resources, audit and accounting services. Personnel-related costs consist of salaries, benefits and stock-based compensation. Facilities costs consist of rent and maintenance of facilities. We expect our general and administrative expenses to increase for the foreseeable future due to anticipated increases in headcount to advance our product candidates and as a result of operating as a public company, including expenses related to compliance with the rules and regulations of the SEC, NYSE, additional insurance expenses, investor relations activities and other administrative and professional services. Other Income (Expense), Net Other income (expense) consists of change in fair value of warrant liabilities, interest earned on our cash equivalents and investments, advisory expense related to our investments and realized gains and losses on marketable securities. Æ 24 Æ Results of Operations for The Three and Six Months Ended June 30, 2024 and 2023 Revenues The Company's revenue related to its out-licensing collaborative agreements consist of upfront license fees and research and development services revenue from its collaboration agreements. The following table summarizes total revenue recognized for the three and six months ended June 30, 2024: Æ Æ Three Months Ended June 30, Æ Æ Six Months Ended June 30, Æ Æ 2024 Æ Æ 2024 Æ Revenue Æ Æ Æ Æ Æ Research and development service revenue Æ Æ 1,435 Æ Æ 1,435 Æ Total Æ Æ \$ 1,435 Æ Æ 1,435 Æ Æ Æ Æ Æ There was no revenue in 2023 since the Acquisition of AnHeart completed in

the second quarter of 2024. The following table presents a breakdown of our operating expenses by functional category:

	Three Months Ended June 30, 2024	Six Months Ended June 30, 2024
Research and development	\$29,247	\$18,590
General and administrative	\$10,657	\$42,089
Operating expenses	\$37,377	\$4,712
Acquired in-process research and development	\$425,070	\$425,070
General and administrative	\$16,156	\$7,541
Research and Development Expenses	\$470,473	\$26,131
Research and Development Expenses	\$444,342	\$490,672
Research and Development Expenses	\$52,652	\$438,020

development expenses increased by \$10.6 million for the three months ended June 30, 2024 compared to 2023. The increase was primarily due to a \$5.9 million increase in personnel-related costs driven by the Acquisition of AnHeart, stock-based compensation and other benefits and a \$4.7 million increase in third-party costs related to research services and drug manufacturing as a result of clinical trial expense for taletrectinib. Research and development expenses increased by \$4.7 million for the six months ended June 30, 2024 compared to 2023. The increase was primarily due to a \$6.6 million increase in personnel-related costs driven by the Acquisition of AnHeart as well as stock-based compensation and other benefits offset by \$1.9 million decrease in third-party costs related to research services and drug manufacturing as a result of completing the Phase 1 monotherapy study of NUV-868 . Acquired In-process Research and Development Expenses On April 9, 2024, as a result of the Acquisition of AnHeart, we recorded a \$425.1 million charge representing an acquired in-process research and development asset with no alternative future use in acquired in-process research and development expenses. General and Administrative Expenses General and administrative expenses increased by \$8.6 million for the three months ended June 30, 2024, compared to 2023. The increase was due to a \$3.9 million increase in personnel-related costs as a result of the Acquisition of AnHeart, \$1.1 million increase in professional fees, \$1.2 million increase in marketing expense, \$0.8 million increase in legal fees, \$0.2 million increase in occupancy expense, \$0.2 million increase in foreign currency impact and a \$1.4 million increase in miscellaneous expense offset by a \$0.2 million decrease in insurance expense. General and administrative expenses increased by \$8.2 million for the six months ended June 30, 2024, compared to 2023. The increase was due to a \$3.8 million increase in personnel-related costs as a result of the Acquisition of AnHeart, \$1.2 million increase in professional fees, \$1.2 million increase in marketing expense, \$0.6 million increase in legal fees, \$0.3 million increase in occupancy expense, \$0.2 million in foreign currency impact and a \$1.5 million increase in miscellaneous expense offset by a \$0.6 million decrease in insurance. Other Income (Expense), Net The following table presents a breakdown of our other income (expense):

	Three Months Ended June 30, 2024	Six Months Ended June 30, 2024
Other income (expense)	\$7,144	\$6,086
Interest income	\$1,058	\$14,274
Interest expense	(132)	(132)
Investment advisory fees	(247)	(231)
Change in fair value of warrant liability	1,135	1,400
Realized loss on marketable securities	(7)	(99)
Total other income (expense), net	\$7,893	\$5,491

Realized loss on marketable securities \$1,300 Other income (expense), net increased by \$2.4 million for the three months ended June 30, 2024 compared to 2023 primarily related to interest income from investments increase of \$1.0 million in 2024 primarily because of higher treasury yield in 2024 compared to 2023, a \$1.4 million increase in the change of fair value of warrant compared to prior year and a decrease of net loss of \$0.1 million on marketable securities offset by a \$0.1 million increase in interest expense. Other income (expense), net increased by \$3.0 million for the six months ended June 30, 2024 compared to 2023 primarily related to interest income from investments increase of \$3.2 million in 2024 primarily because of higher treasury yield in 2024 compared to 2023 and a decrease of net loss of \$0.2 million was primarily due to sale of U.S. Treasuries at losses in 2023 offset by a \$0.2 million increase in the change of fair value of warrant compared to prior year, \$0.1 million increase in interest expense and a \$0.1 million increase in investment advisory fees. Liquidity, Capital Resources and Plan of Operations From inception through June 30, 2024, our operations have been financed primarily by the sale and issuance of Series A preferred stock and Common Stock. As of June 30, 2024, we had \$577.2 million in cash, cash equivalents and marketable securities and an accumulated deficit of \$820.1 million. Our primary use of cash is to fund operating expenses, which consist of research and development expenses related to our clinical-stage product candidates and preclinical programs, and to a lesser extent, general and administrative expenses. Cash used to fund operating expenses is impacted by the timing of when we pay these expenses, as reflected in the change in our outstanding accounts payable and accrued expenses. Based upon our current operating plan, we believe that our existing cash, cash equivalents and marketable securities as of June 30, 2024, will enable us to fund our operating expenses and capital expenditure requirements through at least the next 12 months. We expect to incur substantial expenses in the foreseeable future for the development and potential commercialization of our product candidates and ongoing internal research and development programs. At this time, we cannot reasonably estimate the nature, timing or aggregate amount of costs for our development, potential commercialization, and internal research and development programs. However, in order to complete our current and future preclinical studies and clinical trials, and to complete the process of obtaining regulatory approval for our product candidates, as well as to build the sales, marketing and distribution infrastructure that we believe will be necessary to commercialize our product candidates, if approved, we will require substantial additional funding in the future. Cash Flows The following table summarizes our cash flows for the periods indicated:

	Six Months Ended June 30, 2024	(In thousands)
Cash used in operating activities	\$(53,026)	\$(36,815)
Cash provided by (used in) investing activities	\$41,815	\$(14,650)
Cash provided by financing activities	\$2,769	\$431
Net increase (decrease) in cash and cash equivalents	\$(8,442)	\$(51,034)

Operating Activities In 2024, cash used in operating activities of \$53.0 million was attributable to a net loss of \$477.3 million, a net change of \$10.7 million in our net operating assets and liabilities offset by non-cash charges of \$435.0 million. The change in operating assets and liabilities was primarily due to a \$4.0 decrease in accrued expenses, \$2.8 million decrease in accounts payable, \$1.7 million increase in prepaid expenses and other current assets, \$1.4 million decrease in contract liabilities, \$0.2 million increase in interest receivable on marketable securities, \$0.5 million increase in other non-current assets and \$0.1 million increase in accounts receivable. The non-cash charges consisted primarily of \$425.1 million of acquired in-process research and development, stock-based compensation of \$14.6 million and changes in fair value of warrant liability of \$0.3 million, \$0.3 million of depreciation and amortization offset by amortization of premium on marketable securities of \$5.2 million and \$0.1 million of non cash lease expense. In 2023, cash used in operating activities of \$36.8 million was attributable to a net loss of \$42.4 million offset by a net change of \$0.2 million in our net operating assets and liabilities and non-cash charges of \$5.4 million. The change in operating assets and liabilities was primarily due to a \$0.5 million increase in accounts payable, \$0.3 million decrease in prepaid expenses offset by \$0.3 million increase in interest receivable on marketable securities and \$0.3 million decrease in accrued expenses. The non-cash charges consisted primarily of stock-based compensation of \$10.4 million, realized loss on marketable securities of \$0.2 million, depreciation and amortization of \$0.1 million and changes in fair value of warrant liability of \$0.1 million offset by amortization of premium on marketable securities of \$5.4 million. Investing Activities In 2024, cash provided by investing activities of \$41.8 million was related to the \$251.3 million of proceeds from the sale of marketable securities, \$19.8 cash acquired from the Acquisition of AnHeart, offset by purchase of marketable securities of \$221.9 million and \$7.4 million of transaction costs related to the Acquisition of AnHeart. In 2023, cash used for investing activities of \$14.6 million was related to the purchase of marketable securities of \$443.8 million offset by \$429.2 million of proceeds from the sale of marketable securities. Financing Activities In 2024, cash provided by financing activities of \$2.8 million was related to the \$2.0 million proceeds from exercise of options and \$0.1 million of proceeds from issuance of Common Stock under the Employee Stock Purchase Plan, \$7.0 million of proceeds from borrowings offset by \$6.3 million of debt repayments. In 2023, cash provided by financing activities of \$0.4 million was related to the \$0.3 million of proceeds from exercise of options and \$0.1 million of proceeds from issuance of Common Stock under the Employee Stock Purchase Plan. Critical Accounting Policies and Significant Judgments and Estimates Our management's discussion and analysis of financial condition and results of operations is based upon our consolidated financial statements, which have been prepared in accordance with GAAP. The preparation of these consolidated financial statements requires us to make estimates and judgments that affect the reported amounts of assets, liabilities, revenues and expenses. On an on-going basis, we evaluate our critical accounting policies and estimates. We base our estimates on historical experience and on various other assumptions that we believe to be reasonable in the circumstances, the results of which form the basis for making judgments about the carrying values of assets and liabilities that are not readily apparent from other sources. Actual results may differ from these estimates under different assumptions and conditions. While our significant accounting policies are described in the notes to our consolidated financial statements, we believe that the following critical accounting policies are most important to understanding and evaluating our reported financial results. Revenue Recognition The Company applies ASC Topic 606, "Revenue from Contracts with Customers" ("ASC 606") to account for its revenue transactions. Under ASC 606, an entity recognizes revenue when its customer obtains control of promised goods or services, in an amount that reflects the consideration that the entity expects to receive in exchange for those goods or services. To determine revenue recognition for arrangements that an entity determines are within the scope of ASC 606, the entity performs the five-step model. The Company only applies the five-step model to contracts when it is probable that the entity will collect the consideration to which it is entitled in exchange for the goods or services it transfers to the customer. Once a contract is determined to be within the scope of ASC 606 at contract inception, the Company reviews the contract to determine which performance obligations it must deliver and which of these performance obligations are distinct. The Company recognizes as revenue the amount of the transaction price that is allocated to each performance obligation when that performance obligation is satisfied or as it is satisfied. Accounts receivable represent amounts invoiced and revenues recognized prior to invoicing when the Company has satisfied its performance obligation and has the unconditional right to payment. Collaborative Arrangement In November 2018, the FASB issued Topic 808, "Collaborative Arrangements" ("ASC 808"): Clarifying the Interaction between ASC 808 and ASC 606. This update clarifies that certain transactions between participants in a collaborative arrangement at 27 A should be accounted for under ASC 606 when the counterparty is a customer and precludes an entity from presenting consideration from a transaction in a collaborative arrangement as revenue from contracts with customers if the counterparty is not a customer for that transaction. The Company adopted this standard for all periods presented. The Company enters into collaborative arrangements for the research and development, manufacture and/or commercialization of drug products and drug candidates. The Company assessed and determined that none of the collaboration agreements entered into during the periods presented were within the scope of ASC 808, as all of the agreements did not involve active participation by both parties in a joint research activity, and therefore did not qualify as collaborative arrangements under ASC 808. The Company has determined that all the elements of the above collaborations are reflective of a vendor-customer relationship and therefore within the scope of ASC 606. In determining the appropriate amount of revenue to be recognized as it fulfills its obligations under each of its agreements, the Company performs the five-step model under ASC 606 noted above. The Company recognizes revenue to depict the transfer of control of promised goods or services to customers in an amount that reflects the consideration to which the entity expects to receive in exchange for those goods or services. The Company's collaborative arrangements may contain more than one unit of account, or performance obligation, including grants of licenses to intellectual property rights, agreements to provide research and development services and other deliverables. The collaborative arrangements do not include a right of return for any deliverable. As part of the accounting for these arrangements, the Company must develop assumptions that require judgment to determine the stand-alone selling price for each performance obligation identified in the contract. In developing the stand-alone selling price for a performance obligation, the Company considers competitor pricing for a similar or identical product, market awareness of and perception of the product, expected product life and current market trends. In general, the consideration allocated to each performance obligation is recognized when the respective obligation is satisfied either by delivering a good or providing a service, limited to the consideration that is not constrained. Non-refundable payments received before all of the relevant criteria for revenue recognition are satisfied are recorded as contract liabilities. Licenses of intellectual property If a license to the Company's intellectual property is determined to be distinct from the other promises or performance obligations identified in the contract, the Company recognizes revenue from the portion of the transaction price allocated to the license at a point in time, when the license is transferred to the customer and the customer is able to benefit from the license. Research and development services The portion of a transaction price allocated to research and development services performance obligations is deferred and recognized as revenue over time as delivery or performance of such services occurs based on the use of an input method. Customer options A customer's right to choose, at its discretion, to make a payment for additional goods or services is generally considered an option. If the Company is not presently obligated to provide and does not have a right to consideration for delivering additional goods or services, the item is considered an option. The Company evaluates the customer options for material rights, such as the ability to acquire additional goods or services for free or a discount. Optional future services that reflect their standalone selling prices do not provide the customer with a material right and, therefore, are not considered performance obligations and are accounted for as separate contracts. The optional future services do not include a material right to be accounted for as performance obligations. Milestones payments The Company's collaboration agreements include development and regulatory milestones. The Company evaluates whether the milestones are considered probable of being reached and estimates the amount to be included in the transaction price using the most likely amount method. The Company evaluates factors such as the scientific, clinical, regulatory, commercial, and other risks that must be overcome to achieve the particular milestone in making this assessment. If it is probable that a significant revenue reversal would not occur, the associated milestone value is included in the transaction price. Milestone payments that are not within the Company's control or the licensee's control, such as regulatory approvals, are not considered probable of being achieved until those approvals are received. At the end of each reporting period, the Company re-evaluates the probability of achievement of such milestones and any related constraint, and if necessary, adjusts the estimate of the overall transaction price. Any such adjustments are recorded on a cumulative catch-up basis, which would affect revenue and net loss in the period of adjustment. Royalties For sales-based royalties, including milestone payments based on the level of sales, the Company determines whether the sole or predominant item to which the royalties relate is a license. When the license is the sole or predominant item to which the sales-based royalty relates, the Company recognizes revenue at the later of: (i) when the related sales occur, or (ii) when the performance obligation to which some or all of the royalty has been allocated has been satisfied (or partially satisfied). To date, the Company has not recognized any sales-based royalty revenue resulting from the Company's collaboration agreements. A 28 A The Company receives payments from its customers based on billing terms established in the contract. Up-front payments and fees are recorded as contract liabilities (e.g., deferred revenue) upon receipt or when due until the Company performs its obligations under the arrangement. In the event of an early termination of a collaboration agreement, any contract liabilities would be recognized in the period in which all our obligations under the agreement have been fulfilled. For a complete discussion of accounting for revenue, see the section titled "Collaboration and License Agreements" in Note 5 to our consolidated financial statements for the three months ended June 30, 2024 appearing elsewhere in this report. Research and Development Expenses We expense all research and development costs in the periods in which they are incurred. Costs for certain development activities are recognized based on an evaluation of the progress to completion of specific tasks and estimates of services performed using information and data provided to us by our vendors and third-party service providers. Nonrefundable advance payments for goods or services to be received in future periods for use in research and development activities are deferred and capitalized. The capitalized amounts are then expensed as the related goods are delivered and as services are performed. We expense in-process research and development projects acquired as part of asset acquisitions that have no alternative future use. Warrant Liability We account for warrants as either equity-classified or liability-classified instruments based on an assessment of the warrant's specific terms and applicable authoritative guidance in Financial Accounting Standards Board (FASB) Accounting Standards Codification (ASC) 480, Distinguishing Liabilities from Equity (ASC 480) and ASC 815, Derivatives and Hedging (ASC 815). The assessment considers whether the warrants are freestanding financial instruments pursuant to ASC 480, meet the definition of a liability pursuant to ASC 480, and whether the warrants meet all of the requirements for equity classification under ASC 815, including whether the warrants are indexed to the company's own ordinary shares, among other conditions for equity classification. This assessment, which requires the use of professional judgment, is conducted at the time of warrant issuance and as of each subsequent quarterly period end date while the warrants are outstanding. For issued or modified warrants that meet all of the criteria for equity classification, the warrants are required to be recorded as a component of additional paid-in capital at the time of issuance. For issued or modified warrants that do not meet all the criteria for equity classification, the warrants are required to be recorded as a liability at their fair value on the date of issuance and each balance sheet date thereafter. Changes in the estimated fair value of the warrants are recognized as a non-cash gain or loss on the statements of operations. The fair value of Consideration, Public, and Forward Purchase Warrants was determined using the closing price of the publicly traded warrants on the NYSE market. The fair value of the Private Warrants was estimated using the Black-Scholes option pricing formula. See the section titled "Fair Value

Measurements and Marketable Securities Available-for-Sale" in Note 4 to our consolidated financial statements for the three months ended June 30, 2024 appearing elsewhere in this report. Stock-Based Compensation Expense We estimate the fair value of our stock-based awards to employees and non-employees that are based on a service condition only using the Black-Scholes option-pricing model, which is impacted by our Common Stock price as well as other variables including, but not limited to, expected term that options will remain outstanding, expected Common Stock price volatility over the term of the option awards, risk-free interest rates and expected dividends. We determine the fair value of stock-based awards that are based on both a service condition and achievement of the first to occur of a market or performance condition using a Monte Carlo simulation. The fair value of a stock-based award is recognized over the period during which a recipient is required to provide services in exchange for the award, known as the requisite service period (usually the vesting period) on a straight-line basis. Stock-based compensation expense is recognized based on the fair value determined on the date of grant and is reduced for forfeitures as they occur. Estimating the fair value of stock-based awards as of the grant date using valuation models, such as the Black-Scholes option pricing model, is affected by assumptions regarding a number of variables. Changes in the assumptions can materially affect the fair value and ultimately how much stock-based compensation expense is recognized. These inputs are subjective and generally require significant analysis and judgment to develop. Expected Termâ€œWe have opted to use the â€œsimplified methodâ€œ for estimating the expected term of options whose vesting is based on service condition only, whereby the expected term equals the arithmetic average of the vesting term and the original contractual term of the option (generally 10 years). Expected Volatilityâ€œDue to our limited operating history and a lack of company specific historical and implied volatility data, we have based our estimate of expected volatility on the historical volatility of a group of similar companies that are publicly traded. A 29 % Risk-Free Interest Rateâ€œThe risk-free rate assumption is based on the U.S. Treasury instruments with maturities similar to the expected term of our stock options at the time of the grant. Expected Dividendâ€œWe have not issued any dividends in our history and do not expect to issue dividends over the life of the options and therefore have estimated the dividend yield to be zero. We will continue to use judgment in evaluating the expected volatility, and interest rates utilized for our stock-based compensation expense calculations on a prospective basis. Recent Accounting Pronouncements For information about recent accounting pronouncements, see the sections titled â€œSignificant Accounting Policiesâ€œRecent Accounting Pronouncementsâ€œ in Note 2 to our consolidated financial statements for the three months ended June 30, 2024 appearing elsewhere in this report. Item 3. Quantitative and Qualitative Disclosures About Market Risk. Interest Rate Risk We had cash and investments of \$577.2 million as of June 30, 2024, consisting of cash, money market funds, government securities, and corporate bonds. To date, fluctuations in interest income have not been significant. We do not enter into investments for trading or speculative purposes and have not used any derivative financial instruments to manage our interest rate risk exposure. We have not been exposed to, nor do we anticipate being exposed to, material risks due to changes in interest rates. Foreign Currency Risk Our expenses are generally denominated in U.S. dollars. A 10% increase or decrease in current exchange rates would not have a material effect on our financial results. Item 4. Controls and Procedures. Disclosure Controls and Procedures Our disclosure controls and procedures are designed to ensure that information required to be disclosed in the reports that we file or submit under the Exchange Act is recorded, processed, summarized and reported within the time periods specified in the rules and forms of the Securities and Exchange Commission, and that such information is accumulated and communicated to our management, including our Chief Executive Officer and Principal Financial and Accounting Officer, as appropriate, to allow timely decisions regarding required disclosure. Our management, with the participation of our Chief Executive Officer and Principal Financial and Accounting Officer, evaluated the effectiveness of our disclosure controls and procedures (as defined in Exchange Act Rule 13aâ€œ(15)(e) and 15d-15)(e)) as of June 30, 2024. Based on that evaluation, the Chief Executive Officer and Principal Financial and Accounting Officer concluded that the Companyâ€œs disclosure controls and procedures were effective as of June 30, 2024. Changes in Internal Control Over Financial Reporting On April 9, 2024, we completed the Acquisition of AnHeart Therapeutics Ltd. (â€œAnHeartâ€œ), and are currently integrating AnHeart into our operations, compliance programs and internal control processes. Other than the addition of AnHeartâ€œs operations to our internal control over financial reporting and any related changes in control to integrate AnHeart into Nuvation Bio Inc., there has not been any change in the Companyâ€œs internal control over financial reporting (as such term is defined in Rules 13a-15(f) and 15d-15(f) under the Exchange Act) that has materially affected, or is reasonably likely to materially affect, the Companyâ€œs internal control over financial reporting. A 30 % PART IIâ€œOTHER INFORMATION Item 1. Legal Proceedings. None. Item 1A. Risk Factors. Summary of Risk Factors Below is a summary of material factors that make an investment in our securities speculative or risky. Importantly, this summary does not address all of the risks and uncertainties that we face. Additional discussion of the risks and uncertainties summarized in this risk factor summary, as well as other risks and uncertainties that we face, can be found in the section below. The below summary is qualified in its entirety by that more complete discussion of such risks and uncertainties. You should consider carefully the risks and uncertainties described below as part of your evaluation of an investment in our securities: â€œWe have a limited operating history and have incurred significant losses since inception and anticipate that we may continue to incur losses for the foreseeable future, and may never achieve or maintain profitability. â€œWe will need substantial funding to pursue our business objectives. If we are unable to raise capital when needed or on favorable terms, we could be forced to delay, reduce or terminate our product development, other operations or commercialization efforts. Additionally, raising additional capital may cause dilution to our stockholders, restrict our operations or require us to relinquish proprietary rights. â€œFollowing our recent Acquisition of AnHeart, we may have difficulty integrating AnHeartâ€œs business, operations and employees, including those based in China. This may cause disruption to ongoing business, divert the attention of management, and the process of combining the companies may cost more and take longer than originally anticipated. â€œIf we do not obtain regulatory approval for and successfully commercialize our product candidates in one or more indications or we experience significant delays in doing so, we may never generate any revenue or become profitable. â€œOur approach to the discovery and development of product candidates based on our Drug-Drug Conjugate platform is unproven and is based on novel technology, and we do not know whether we will be able to develop any products of commercial value, or if competing technological approaches will limit the commercial value of our product candidates or render our platform obsolete. â€œClinical trials are very expensive, time-consuming and difficult to design and implement, and involve uncertain outcomes. Furthermore, results of earlier preclinical studies and clinical trials may not be predictive of results of future preclinical studies or clinical trials. â€œWe may encounter substantial delays in our preclinical studies or clinical trials or we may fail to demonstrate safety and efficacy to the satisfaction of applicable regulatory authorities. â€œIf any of our product candidates receives marketing approval and we, or others, later discover that the drug is less effective than previously believed or causes undesirable side effects that were not previously identified, our ability to market the drug could be compromised. â€œWe may become exposed to costly and damaging liability claims, either when testing our product candidates in the clinic or at the commercial stage, and our product liability insurance may not cover all damages from such claims. â€œWe have never commercialized a product candidate and we may lack the necessary expertise, personnel and resources to successfully commercialize any of our products that receive regulatory approval on our own or together with collaborators. â€œWe face substantial competition, which may result in others discovering, developing or commercializing products before or more successfully than we do. â€œEven if we obtain regulatory approval for our product candidates, they will remain subject to ongoing regulatory oversight. â€œWe rely on third parties to perform the chemistry work associated with our drug discovery and preclinical activities and to conduct our preclinical studies and future clinical trials, and our business could be substantially harmed if these third parties cease performing services or perform in an unsatisfactory manner. â€œWe do not have our own manufacturing capabilities and will rely on third parties to produce clinical and commercial supplies of taltirecinib, safusidenib, NUV-868, NUV-1511 and our other current and future product candidates. A 31 % â€œIf we are not able to maintain existing collaborations or establish new collaborations, we may have to alter some of our future development and commercialization plans. â€œOur business operations and current and future relationships with investigators, healthcare professionals, consultants, third-party payors and customers will be subject, directly or indirectly, to federal and state healthcare fraud and abuse laws, false claims laws, transparency laws, health information privacy and security laws and other healthcare laws and regulations including equivalent foreign laws and regulations. If we are unable to comply, or have not fully complied, with such laws, we could face substantial penalties. â€œIf we are unable to obtain, maintain, protect and enforce sufficient patent and other intellectual property rights for our product candidates and technology, or if the scope of patent and other intellectual property rights obtained is not sufficiently broad, we may not be able to compete effectively in our market. â€œOur reliance on third parties requires us to share our trade secrets, which increases the possibility that a competitor will discover them or that our trade secrets will be misappropriated or disclosed. â€œWe may become involved in lawsuits to protect or enforce our patents or other intellectual property, which could be expensive, time-consuming and unsuccessful, and issued patents covering our technology and product candidates could be found invalid or unenforceable if challenged. â€œThird parties may initiate legal proceedings alleging that we are infringing, misappropriating or otherwise violating their intellectual property rights, the outcome of which would be uncertain and could negatively impact the success of our business. â€œIntellectual property litigation could cause us to spend substantial resources and distract our personnel from their normal responsibilities. â€œOur business, operations and clinical development plans and timelines and supply chain could be adversely affected by the effects of health epidemics, on the manufacturing, clinical trial and other business activities performed by us or by third parties with whom we conduct business, including our contract manufacturing organizations (â€œCMOsâ€œ), contract research organizations (â€œCROsâ€œ), shippers and others. â€œOur future success depends on our ability to retain Dr. Hung and our other key employees, consultants and advisors and to attract, retain and motivate qualified personnel. â€œThe dual-class structure of our Common Stock has the effect of concentrating voting power with our Chief Executive Officer, which limits other stockholdersâ€œ ability to influence the outcome of important transactions, including a change in control. Risks Related to Our Financial Position and Need for Additional Capital We have a limited operating history and have incurred significant losses since inception and anticipate that we may continue to incur losses for the foreseeable future, and may never achieve or maintain profitability. Biopharmaceutical product development is a highly speculative undertaking and involves a substantial degree of risk. We are an oncology company with a limited operating history upon which you can evaluate our business and prospects. We commenced operations in 2018, and to date, we have focused primarily on organizing and staffing our company, business planning, raising capital, identifying product candidates, establishing our intellectual property portfolio and conducting research, preclinical studies and clinical trials. Our approach to the discovery and development of product candidates is unproven, and we do not know whether we will be able to develop any product candidates that succeed in clinical development or products of commercial value. As an organization, we have not yet completed any clinical trials, obtained regulatory approvals, manufactured a commercial-scale product (or arranged for a third party to do so on our behalf), or conducted sales and marketing activities necessary for successful product commercialization. Consequently, any predictions made about our future success or viability may not be as accurate as they could be if we had a history of successfully developing and commercializing biopharmaceutical products. Since inception, we have not generated any product revenue and have incurred significant operating losses. Our net losses were \$104.2 million and \$75.8 million in 2022 and 2023, respectively. As of June 30, 2024, we had an accumulated deficit of \$820.1 million. We expect to continue to incur significant expenses and increasing operating losses for the foreseeable future. Since inception, we have devoted substantially all of our efforts to research and preclinical and clinical development of our product candidates, as well as to building our management team and infrastructure. It could be at least several years, if ever, before we have a commercialized drug. The net losses we incur may fluctuate significantly from quarter to quarter and year to year. We anticipate that our expenses will increase substantially if, and as, we: â€œcontinue to advance our research and preclinical and clinical development of our product candidates; A 32 % â€œexpand and initiate further clinical trials for our product candidates; â€œseek to identify additional product candidates; â€œseek marketing approvals for our product candidates that successfully complete clinical trials, if any; â€œestablish a sales, marketing and distribution infrastructure to commercialize any products for which we may obtain marketing approval; â€œmaintain, expand, protect and enforce our intellectual property portfolio and obtain licenses to third-party intellectual property; â€œattract, hire and retain additional administrative, clinical, regulatory and scientific personnel; â€œenter into third-party relationships for clinical trials, manufacturing and supply; and â€œincur additional legal, accounting and other expenses in operating our business, including the additional costs associated with operating as a public company. In addition, because of the numerous risks and uncertainties associated with pharmaceutical products and development, we are unable to accurately predict the timing or amount of increased expenses and when, or if, we will be able to achieve profitability. Our expenses could increase and profitability could be further delayed if we decide to or are required by the FDA or other comparable foreign regulatory authorities such as the European Medicines Agency (â€œEMAâ€œ), the U.K. Medicines & Healthcare Products Regulatory Agency (the â€œMHRAâ€œ), or the National Medical Product Administration of China (the â€œNMPAâ€œ), to perform studies or trials in addition to those currently expected, or if there are any delays in the development or completion of any current or future preclinical studies or clinical trials of our current and future product candidates. Even if we complete the development and regulatory processes described above, we anticipate incurring significant costs associated with launching and commercializing our current and future product candidates. Even if we do achieve profitability, we may not be able to sustain or increase profitability on a quarterly or annual basis. Our failure to become and remain profitable would decrease our value and could impair our ability to raise capital, maintain our research and development efforts, expand our business or continue our operations. A decline in value also could cause you to lose all or part of your investment. We will need substantial funding to pursue our business objectives. If we are unable to raise capital when needed or on favorable terms, we could be forced to delay, reduce or terminate our product development, other operations or commercialization efforts. Identifying and developing potential product candidates and conducting preclinical studies and clinical trials is a time-consuming, expensive and uncertain process that takes years to complete, and we may never generate the necessary data or results required to obtain regulatory approval and begin selling any approved products. We expect our expenses to increase in connection with our ongoing activities, particularly as we conduct our ongoing and planned preclinical studies and clinical trials, initiate additional clinical trials for our product candidates and seek regulatory approval for our current product candidates and any future product candidates we may develop. Our expenses could increase beyond our current expectations if the FDA, or comparable foreign regulatory authorities, require us to perform clinical trials and other studies in addition to those that we currently anticipate. In addition, if we obtain marketing approval for any of our product candidates, we expect to incur significant commercialization expenses related to product sales, marketing, manufacturing and distribution. Accordingly, we will need to obtain substantial additional funding in connection with our continuing operations. If we are unable to raise capital when needed or on attractive terms, we will be forced to delay, reduce or terminate our research and development programs or future commercialization efforts. As of June 30, 2024, we had \$577.2 million in cash and investments, and an accumulated deficit of \$820.1 million. Based upon our current operating plan, we believe that our existing cash, cash equivalents and marketable securities will be sufficient to fund our operations for at least the next 12 months. This estimate is based on assumptions that may prove to be wrong, and we could use our available capital resources sooner than we expect. Changes may occur beyond our control that would cause us to consume our available capital before that time, including changes in and progress of our development activities and changes in regulation. Our future capital requirements will depend on many factors, including: â€œthe scope, rate of progress, results and costs of drug discovery, preclinical development, laboratory testing and clinical trials for our product candidates; â€œthe number and development requirements of product candidates that we may pursue, and other indications for our current product candidates that we may pursue; â€œthe costs, timing and outcome of regulatory review of our product candidates; â€œthe scope and costs of manufacturing development and commercial manufacturing activities; â€œthe cost associated with commercializing any approved product candidates; A 33 % â€œthe cost and timing of developing our ability to establish sales and marketing capabilities, if any; â€œthe costs of preparing, filing and prosecuting patent applications, maintaining, enforcing and protecting our intellectual property rights, defending intellectual property-related claims and obtaining licenses to third-party intellectual property; â€œour ability to establish and maintain collaborations on favorable terms, if at all; and â€œthe extent to which we acquire or in-license other product candidates and technologies and associated intellectual property. We will require additional capital to complete our planned clinical development programs for our clinical stage product candidates as well as our preclinical product candidates to obtain regulatory approval. Any additional capital raising efforts may divert our management from their day-to-day activities, which may adversely affect our ability to develop and commercialize our current and future product candidates, if approved. In addition, we cannot guarantee that future financing will be available on a timely basis, in sufficient amounts or on terms acceptable to us, if at all. Moreover, the terms of any financing may adversely affect the holdings or the rights of our stockholders and our issuance of additional securities, whether equity or debt, or the market perception that such issuances are likely to occur, could cause the market price of our Common Stock to decline. If we are unable to obtain funding on a timely basis on acceptable terms, we may be required to delay, reduce or terminate one or more of our research and development programs or the commercialization of any product candidates that may be approved. This could harm our business and could potentially cause us to cease operations. Raising additional capital may cause dilution to our stockholders, restrict our operations or require us to relinquish proprietary rights. Until such time, if ever, as we can generate substantial product revenues, we expect to finance

our cash needs through a combination of equity offerings, debt financings, collaborations, strategic alliances and licensing arrangements. To the extent that we raise additional capital through the sale of equity or convertible debt securities, your ownership interest will be diluted and the terms of these securities may include liquidation or other preferences that adversely affect your rights as a stockholder. Debt financing, if available, may involve agreements that include covenants limiting or restricting our ability to take specific actions, such as incurring additional debt, making capital expenditures or declaring dividends. If we raise additional funds through collaborations, strategic alliances or licensing arrangements with third parties, we may have to relinquish valuable rights to our technologies, future revenue streams, research programs or product candidates or grant licenses on terms that may not be favorable to us. If we are unable to raise additional funds through equity or debt financings when needed, we may be required to delay, reduce or terminate our product development or future commercialization efforts or grant rights to third parties to develop and market product candidates that we would otherwise prefer to develop and market ourselves. Changes in tax laws or regulations that are applied adversely to us or our customers may have a material adverse effect on our business, cash flow, financial condition or results of operations. New income, sales, use or other tax laws, statutes, rules, regulations or ordinances could be enacted at any time, which could affect the tax treatment of our earnings. Any new taxes could adversely affect our business operations, and our business and financial performance. Further, existing tax laws, statutes, rules, regulations or ordinances could be interpreted, changed, modified or applied adversely to us, which could require us to pay additional taxes on a prospective or retroactive basis, as well as penalties, interest and other costs, including compliance costs. The Tax Cuts and Jobs Act enacted in 2017 (the “Tax Act”), the Coronavirus Aid, Relief, and Economic Security Act enacted in 2020 and the Inflation Reduction Act enacted in 2022 made many significant changes to the U.S. tax laws. For example, effective January 1, 2022, the Tax Act eliminated the option to deduct research and development expenses for tax purposes in the year incurred and instead requires taxpayers to capitalize and subsequently amortize such expenses over five years for research activities conducted in the United States and over 15 years for research activities conducted outside the United States. Although there have been legislative proposals to repeal or defer the capitalization requirement to later years, there can be no assurance that the provision will be repealed or otherwise modified. Future guidance from the Internal Revenue Service and other tax authorities with respect to any such legislation may affect us, and certain aspects of such legislation could be repealed or modified in future legislation. In addition, it is uncertain if and to what extent various states will conform to federal tax laws. Future tax reform legislation could have a material impact on the value of our deferred tax assets, could result in significant one-time charges, and could increase our future U.S. tax expense. In addition, our tax obligations and effective tax rate could increase, including as a result of the base erosion and profit shifting (“BEPS”) project that is being led by the Organization for Economic Co-operation and Development (“OECD”), and other initiatives led by the OECD. For example, the OECD is leading work on proposals, commonly referred to as BEPS 2.0, which, if and to the extent implemented, would make important changes to the international tax system. These proposals are based on two “pillars”, involving the reallocation of taxing rights in respect of certain multinational enterprises above a fixed profit margin to the jurisdictions in which they carry on business (referred to as “Pillar One”) and imposing a minimum effective corporate tax rate on certain “large multinational enterprises (referred to as “Pillar Two”). A number of countries have enacted with effect from January 1, 2024, or are in the process of enacting, core elements of the Pillar Two rules. We are monitoring developments and evaluating the potential impacts of these new rules on our business. Our ability to use our net operating loss carryforwards and certain other tax attributes to offset taxable income or taxes may be limited. As of December 31, 2023, we had federal and state net operating loss (“NOL”) carryforwards of \$95.2 million and \$140.3 million, respectively. Under current law, federal NOLs incurred in taxable years beginning after December 31, 2017 may be carried forward indefinitely, but such federal NOL carryforwards are permitted to be used in any taxable year to offset only up to 80% of taxable income in such year. Separately, under Sections 382 and 383 of the Internal Revenue Code of 1986, as amended, or the Internal Revenue Code, and corresponding provisions of state law, if a corporation undergoes an “ownership change,” which generally is defined as a greater than 50% change, by value, in its equity ownership over a three-year period, the corporation’s ability to use its pre-change NOL carryforwards and other pre-change tax attributes, such as research tax credits, to offset its post-change income or taxes may be limited. The completion of the 2021 merger of Nuvation Bio Inc. and Panacea Acquisition Corp., together with private placements and other transactions that have occurred since our inception, may have triggered such an ownership change pursuant to Section 382. We have not completed a Section 382 analysis, and therefore, there can be no assurances that our NOLs are not already limited. We also may experience ownership changes as a result of subsequent shifts in our stock ownership, some of which may be outside of our control. As a result, if we earn net taxable income, our ability to use our pre-change NOL carryforwards to offset U.S. federal taxable income may be subject to limitations, which potentially could result in increased future tax liability to us. If an ownership change occurs and our ability to use our net operating loss carryforwards is materially limited, this could harm our future operating results by effectively increasing our future tax obligations. In addition, due to changes in laws and regulations, including changes proposed or implemented by the current or a future U.S. presidential administration, such as alternative minimum taxes, or other unforeseen reasons, our existing net operating losses could become unavailable to reduce future income tax liabilities. Further, at the state level, there may be periods during which the use of NOL carryforwards is suspended or otherwise limited, which could accelerate or permanently increase state taxes owed. For example, California imposed limits on the usability of California state net operating losses to offset taxable income in tax years beginning after 2023 and before 2027. Risks Related to the Development of our Product Candidates If we do not obtain regulatory approval for and successfully commercialize our product candidates in one or more indications or we experience significant delays in doing so, we may never generate any revenue or become profitable. We do not have any products that have received regulatory approval and may never be able to develop marketable product candidates. We have invested substantially all of our efforts in developing and identifying potential product candidates and conducting preclinical and clinical studies. As a result, our business currently depends heavily on the successful development, regulatory approval and, if approved, commercialization of taltrectinib, safusidenib, NUV-868 and NUV-1511. We cannot be certain that any of these or any other product candidates will receive regulatory approval or will be successfully commercialized even if we receive regulatory approval. The research, testing, manufacturing, safety, efficacy, labeling, approval, sale, marketing and distribution of product candidates is, and will remain, subject to comprehensive regulation by the FDA and similar foreign regulatory authorities. Before obtaining regulatory approvals for the commercial sale of any product candidate, we must demonstrate through preclinical studies and clinical trials that the product candidate is safe and effective for use in each target indication. Drug development is a long, expensive and uncertain process, and delay or failure can occur at any stage of any of our preclinical studies or clinical trials. For example, based on our preclinical or clinical experience since February 2022, we have discontinued or deprioritized three of the five programs, including the lead program, that we were pursuing at that time. Failure to obtain regulatory approval for our product candidates will prevent us from commercializing and marketing our product candidates. The success of our product candidates will depend on several additional factors, including: successful completion of preclinical studies; successful initiation of clinical trials; successful patient enrollment in, and completion, of clinical trials that demonstrate their safety and efficacy; receiving marketing approvals from applicable regulatory authorities; obtaining, maintaining, protecting and enforcing patent, trade secret and other intellectual property rights and regulatory exclusivity for our product candidates; completing any post-marketing studies required by applicable regulatory authorities; A 35 A making and maintaining arrangements with third-party manufacturers, or establishing manufacturing capabilities, for both clinical and commercial supplies of our product candidates; establishing sales, marketing and distribution capabilities and successfully launching commercial sales of our products, if and when approved, whether alone or in collaboration with others; the prevalence and severity of adverse events experienced with our product candidates; acceptance of our product candidates by patients, the medical community and third-party payors; a continued acceptable safety profile following approval; obtaining and maintaining healthcare coverage and adequate reimbursement for our product candidates; competing effectively with other cancer therapies, including with respect to the sales and marketing of our product candidates, if approved; and obtaining licenses to any third-party intellectual property we deem necessary or desirable. Many of these factors are beyond our control, including the time needed to adequately complete preclinical studies, clinical testing and the regulatory submission process, our ability to obtain and protect intellectual property rights and changes in the competitive landscape. It is possible that none of our product candidates will ever obtain regulatory approval, even if we expend substantial time and resources seeking such approval. If we do not achieve one or more of these factors in a timely manner or at all, we could experience significant delays or an inability to successfully complete clinical trials, obtain regulatory approval or, if approved, commercialize our product candidates, which would materially harm our business, financial condition, results of operations and prospects. In addition, the clinical trial requirements of the FDA, the European Commission, competent authorities of EU Member States, the MHRA, the NMPA and other comparable regulatory authorities and the criteria regulators may use to determine the safety and efficacy of a product candidate vary substantially according to the type, complexity, novelty and intended use and market of the potential products. The regulatory approval process for novel product candidates can be more expensive and take longer than for other, better known or extensively studied pharmaceutical or other product candidates. Our approach to the discovery and development of product candidates based on our DDC platform is unproven, and we do not know whether we will be able to develop any products of commercial value, or if competing technological approaches will limit the commercial value of our product candidates or render our platform obsolete. The success of our business depends in part upon our ability to identify, develop and commercialize products based on our proprietary Drug-Drug Conjugate (“DDC”) platform, which leverages a novel and unproven therapeutic approach within the drug-conjugate class of anti-cancer therapies. While our first DDC clinical product candidate, NUV-1511, is in a Phase 1 dose escalation trial that began in the first quarter of 2024, we have not yet demonstrated safety or efficacy for NUV-1511 or any other DDC product candidate. Our research methodology and novel approach to oncology using our DDC platform may be unsuccessful in identifying additional product candidates, and any product candidates based on our technology may be shown to have harmful side effects or may have other characteristics that may necessitate additional clinical testing, or make the product candidates unmarketable or unlikely to receive marketing approval. In addition, adverse developments with respect to one of our DDC platform-based programs may have a significant adverse impact on the actual or perceived likelihood of success and value of similar programs. In addition, the biotechnology and biopharmaceutical industries are characterized by rapidly advancing technologies. Our future success will depend in part on our ability to maintain a competitive position with our DDC platform. If we fail to stay at the forefront of technological change in utilizing our DDC platform to create and develop product candidates, we may be unable to compete effectively. Our competitors may render our DDC platform obsolete, or limit the commercial value of our product candidates, by advances in existing technological approaches or the development of new or different approaches, potentially eliminating the advantages in our drug discovery process that we believe we derive from our research approach and proprietary technologies. By contrast, adverse developments with respect to other companies that attempt to use a similar approach to our approach may adversely impact the actual or perceived value of our DDC platform and potential of our DDC platform-based product candidates. If any of these events occur, we may be forced to abandon our development efforts for a program or programs, which would harm our business. Our DDC platform-based product candidates are based on a novel technology, which makes it difficult to predict the time and cost of product candidate development. We have concentrated our product research and development efforts on our novel DDC platform, and our future success depends in part on the successful development of product candidates arising from our DDC platform. There can be no assurance that any development problems we may experience in the future related to our DDC platform will not cause significant delays or unanticipated costs, or that such development problems can be efficiently solved. We may also experience delays in developing a sustainable, reproducible and scalable manufacturing process or transferring that process to commercial partners, which may prevent us from completing our clinical trials or commercializing our product candidates on a timely or profitable basis, if at all. We may in the future develop product candidates in combination with other therapies and that may expose us to additional risks. We may develop future product candidates for use in combination with one or more currently approved cancer therapies. Even if any product candidate we develop was to receive marketing approval or be commercialized for use in combination with other existing therapies, we would continue to be subject to the risks that the FDA or similar foreign regulatory authorities could revoke approval of the therapy used in combination with our product candidate or that safety, efficacy, manufacturing or supply issues could arise with these existing therapies. Combination therapies are commonly used for the treatment of cancer, and we would be subject to similar risks if we develop any of our product candidates for use in combination with other drugs or for indications other than cancer. This could result in our own products being removed from the market or being less successful commercially. We may also evaluate our product candidates in combination with one or more other cancer therapies that have not yet been approved for marketing by the FDA or similar foreign regulatory authorities. We will not be able to market and sell our product candidates we develop in combination with any such unapproved cancer therapies that do not ultimately obtain marketing approval. If the FDA or similar foreign regulatory authorities do not approve or revoke the approval of these other drugs, or if safety, efficacy, manufacturing or supply issues arise with the drugs we choose to evaluate in combination with our product candidates, we may be unable to obtain approval of or market our product candidates. Clinical trials are very expensive, time-consuming and difficult to design and implement, and involve uncertain outcomes. Furthermore, results of earlier preclinical studies and clinical trials may not be predictive of results of future preclinical studies or clinical trials. The risk of failure for our product candidates is high. It is impossible to predict when or if any of our product candidates will prove safe or effective in humans or will receive regulatory approval. To obtain the requisite regulatory approvals to market and sell any of our product candidates, we must demonstrate through extensive preclinical studies and clinical trials that our product candidates are safe and effective in humans for use in each target indication. Preclinical investigation and clinical testing is expensive and can take many years to complete, and the outcome is inherently uncertain. Failure can occur at any time during the preclinical investigation or clinical trial process. For example, in August 2022, we announced the discontinuation of development of our former lead program, NUV-422, following the emergence of a safety signal, uveitis, which is a form of inflammation of the eye. In addition, the results of preclinical studies and earlier clinical trials may not be predictive of the results of later-stage preclinical studies or clinical trials. The results generated to date in preclinical studies for our product candidates do not ensure that later preclinical studies or clinical trials will demonstrate similar results. Product candidates in later stages of clinical trials may fail to show the desired safety and efficacy traits despite having progressed through preclinical and earlier stage clinical trials. In later-stage clinical trials, we will likely be subject to more rigorous statistical analyses than in completed earlier stage clinical trials. A number of companies in the pharmaceutical industry have suffered significant setbacks in later-stage clinical trials due to adverse safety profiles or lack of efficacy, notwithstanding promising results in earlier trials, and we cannot be certain that we will not face similar setbacks. Moreover, preclinical and clinical data are often susceptible to varying interpretations and analyses, and many companies that have believed their product candidates performed satisfactorily in preclinical studies and clinical trials have nonetheless failed to obtain marketing approval of their products. In some instances, there can be significant variability in safety or efficacy results between different clinical trials of the same product candidate due to numerous factors, including changes in clinical trial procedures set forth in protocols, differences in the size and type of the patient populations, adherence to the dosing regimen and other clinical trial protocols, and the rate of dropout among clinical trial participants. If we fail to produce positive results in our planned preclinical studies or clinical trials of any of our product candidates, the development timeline and regulatory approval and commercialization prospects for our product candidates, and, correspondingly, our business and financial prospects, would be materially and adversely affected. Interim, topline, and preliminary data from our clinical trials that we announce or publish from time to time may change as more patient data become available and are subject to audit and verification procedures that could result in material changes in the final data. From time to time, we may publicly disclose preliminary or topline data from our preclinical studies and clinical trials, which is based on a preliminary analysis of then-available data, and the results and related findings and conclusions are subject to change following a more comprehensive review of the data related to the particular study or trial. We also make assumptions, estimations, calculations and conclusions as part of our analyses of data, and we may not have received or had the opportunity to fully and carefully evaluate all data. As a result, the topline or preliminary results that we report may differ from future results of the same studies, or different conclusions or considerations may qualify such results, once additional data have been received and fully evaluated. Topline data also remain subject to audit and verification procedures that may result in the final A 37 A data being materially different from the preliminary data we previously published. As a result, topline data should be viewed with caution until the final data are available. From time to time, we may also disclose interim data from our preclinical studies and clinical trials. Interim data from clinical trials that we may complete are subject to the risk that one or more of the clinical outcomes may materially change as patient enrollment continues and more patient data become available. Adverse differences between preliminary or interim data and final data could significantly harm our business prospects. Further, others, including regulatory authorities, may not accept or agree with our assumptions, estimates, calculations, conclusions or analyses or may interpret or weigh the importance of data differently, which could impact the value of the particular program, the approvability or commercialization of the particular drug candidate or

product and our company in general. In addition, the information we choose to publicly disclose regarding a particular study or clinical trial is based on what is typically extensive information, and you or others may not agree with what we determine is material or otherwise appropriate information to include in our disclosure. If the interim, topline, or preliminary data that we report differ from actual results, or if others, including regulatory authorities, disagree with the conclusions reached, our ability to obtain approval for and commercialize our product candidates may be harmed, which could harm our business, operating results, prospects or financial condition. We may encounter substantial delays in our preclinical studies or clinical trials or we may fail to demonstrate safety and efficacy to the satisfaction of applicable regulatory authorities. Before obtaining marketing approval from regulatory authorities for the sale of our product candidates, we must conduct extensive clinical trials to demonstrate the safety and efficacy of the product candidate for its intended indications. Preclinical studies and clinical trials are expensive, time-consuming and uncertain as to outcome. We cannot guarantee that any preclinical studies or clinical trials will be conducted as planned or completed on schedule, if at all. A failure of one or more preclinical studies or clinical trials can occur at any stage of testing. Events that may prevent successful or timely completion of preclinical or clinical development include: delays in conducting experiments or preclinical studies or unsatisfactory results from such experiments or studies; delays in reaching a consensus with regulatory authorities on trial design, dose optimization or dose selection; delays in reaching agreement or failing to agree on acceptable terms with prospective CROs and clinical trial sites; delays in opening sites and recruiting suitable patients to participate in our clinical trials; delays in enrollment due to travel or quarantine policies, or other factors, related to health epidemics, other pandemics or other events outside our control; imposition of a clinical hold by regulatory authorities as a result of a serious adverse event, concerns with a class of product candidates or after an inspection of our clinical trial operations or trial sites; delays in having patients complete participation in a trial or return for post-treatment follow-up; occurrence of serious adverse events associated with the product candidate that are viewed to outweigh its potential benefits; or changes in regulatory requirements and guidance that require amending or submitting new clinical protocols. For instance, health epidemics and the measures taken in response by the governmental authorities could disrupt the supply chain and the manufacture or shipment of drug substances and finished drug products for our product candidates for use in our research and clinical trials, delay, limit or prevent our employees and CROs from continuing research and development activities, impede the ability of patients to enroll or continue in clinical trials, or impede testing, monitoring, data collection and analysis or other related activities, any of which could delay our clinical trials and increase our development costs, and have a material adverse effect on our business, financial condition and results of operations. Any inability to timely and successfully complete preclinical and clinical development could result in additional costs to us or impair our ability to achieve regulatory and commercialization milestones. In addition, if we make manufacturing or formulation changes to our product candidates, we may need to conduct additional testing to bridge our modified product candidate to earlier versions. Clinical trial delays could also shorten any periods during which we may have the exclusive right to commercialize our product candidates, if approved, or allow our competitors to bring comparable drugs to market before we do, which could impair our ability to successfully commercialize our product candidates and may harm our business, financial condition, results of operations and prospects. Additionally, if the results of our clinical trials are inconclusive or if there are safety concerns or serious adverse events associated with our product candidates, we may: A 38 A be delayed or unsuccessful in obtaining marketing approval; obtain approval for indications or patient populations that are not as broad as intended or desired; obtain approval with labeling that includes significant use or distribution restrictions or safety warnings; be subject to additional post-marketing testing requirements; be required to perform additional clinical trials to support approval or be subject to additional post-marketing testing requirements; have regulatory authorities withdraw, vary or suspend, their approval of the drug or impose restrictions on its distribution in the form of a modified risk evaluation and mitigation strategy, or REMS, or comparable foreign restrictions; be subject to the addition of labeling statements, such as warnings or contraindications; be sued; or experience damage to our reputation. Our drug development costs will also increase if we experience delays in testing or obtaining marketing approvals. We do not know whether any of our preclinical studies or clinical trials will begin as planned, need to be restructured or be completed on schedule, if at all. Further, we, the FDA or comparable foreign regulatory authorities, an Institutional Review Board, or an Ethics Committee may suspend our clinical trials at any time if it appears that we or our collaborators are failing to conduct a trial in accordance with applicable regulatory requirements, including the FDA's current Good Clinical Practice, and foreign equivalents, regulators find that we are exposing participants to unacceptable health risks or if the FDA or comparable foreign regulatory authorities find deficiencies in our Investigational New Drugs, clinical trial applications or the conduct of these trials. Therefore, we cannot predict with any certainty the schedule for commencement and completion of future clinical trials. If we experience delays in the commencement or completion of our clinical trials, or if we terminate a clinical trial prior to completion, the commercial prospects of our product candidates could be negatively impacted, and our ability to generate revenues from our product candidates may be delayed or eliminated entirely. If we encounter continued or new difficulties enrolling patients in our clinical trials, our clinical development activities could be delayed or otherwise adversely affected. We have experienced and may in the future experience difficulties in patient enrollment in our clinical trials for a variety of reasons, including challenges resulting from health epidemics, labor shortages, and global supply chain interruptions. The timely completion of clinical trials in accordance with their protocols depends, among other things, on our ability to enroll a sufficient number of patients who remain in the study until its conclusion. The enrollment of patients depends on many factors, including: the patient eligibility criteria defined in the protocol; the size and health of the patient population required for analysis of the trial's primary endpoints; the proximity of patients to study sites; the design of the trial; our ability to recruit clinical trial investigators with the appropriate competencies and experience; clinicians' and patients' perceptions as to the potential advantages of the product candidate being studied in relation to other available therapies, including any new drugs that may be approved for the indications we are investigating; our ability to obtain and maintain patient consents; and the risk that patients enrolled in clinical trials will drop out of the trials before completion. In addition, our clinical trials will compete with other clinical trials for product candidates that are in the same therapeutic areas as our product candidates, and this competition will reduce the number and types of patients available to us, because some patients who might have opted to enroll in our trials may instead opt to enroll in a trial being conducted by one of our competitors. Since the number of qualified clinical investigators is limited, we expect to conduct some of our clinical trials at the same clinical trial sites that some of our competitors use, which will reduce the number of patients who are available for our clinical trials at such clinical trial site. Moreover, because our product candidates represent a departure from more commonly used methods for cancer treatment, potential patients and their doctors may be inclined to use conventional therapies rather than enroll patients in any future clinical trial. A 39 A Delays in patient enrollment may result in increased costs or may affect the timing or outcome of our current or planned clinical trials, which could prevent completion of these trials and adversely affect our ability to advance the development of our product candidates. If we or third parties are unable to successfully develop companion diagnostics for taltretectinib, safusidenib, or any of our other product candidates that are targeted therapies, or if we experience significant delays in doing so, we may not achieve marketing approval or realize the full commercial potential of such product candidates. A key part of our development strategy for taltretectinib, safusidenib, and any of our other product candidates that are targeted therapies, is to identify subsets of patients with specific types of tumors that express specific genetic markers. Identification of these patients may require the development and use of companion diagnostics. The FDA generally will require either approval or clearance of the diagnostic at the same time the FDA approves the therapeutic product, or as a post-marketing commitment at the time of the therapeutic product's approval. We do not have experience or capabilities in developing or commercializing diagnostics and plan to rely in large part on third parties to perform these functions. Companion diagnostics are subject to regulation by the FDA and comparable foreign regulatory authorities as medical devices and may require separate regulatory approval prior to commercialization of the associated product candidate. If we or third parties are unable to successfully develop companion diagnostics for our product candidates, or experience delays in doing so: the development of these product candidates may be delayed because it may be difficult to identify patients for enrollment in our clinical trials in a timely manner; these product candidates may not receive marketing approval if their safe and effective use depends on a companion diagnostic; and we may not realize the full commercial potential of these product candidates that receive marketing approval if, among other reasons, we are unable to appropriately identify patients or types of tumors with the specific genetic alterations targeted by these product candidates. Even if our product candidates and any associated companion diagnostics are approved for marketing, the need for companion diagnostics may slow or limit adoption of our product candidates. Although we believe genetic testing is becoming more prevalent in the diagnosis and treatment of cancer, our product candidates may be perceived negatively compared to alternative treatments that do not require the use of companion diagnostics, due to either the additional cost of the companion diagnostic or the need to complete additional procedures to identify genetic markers prior to administering our product candidates. If any of these events were to occur, our business and growth prospects would be harmed, possibly materially. We may expend our limited resources to pursue a particular product candidate or indication and fail to capitalize on product candidates or indications that may be more profitable or for which there is a greater likelihood of success. Because we have limited financial and managerial resources, we focus on research programs that we identify for specific indications. As a result, we may forego or delay pursuit of opportunities with other product candidates or for other indications, even those that we have begun investigating and that may have shown promise, that later prove to have greater commercial potential. Our resource allocation decisions may cause us to fail to capitalize on viable commercial therapies or profitable market opportunities. Our spending on current and future research and development programs and product candidates for specific indications may not yield any commercially viable products. If we do not accurately evaluate the commercial potential or target market for a particular product candidate, we may relinquish valuable rights to that product candidate through collaboration, licensing or other royalty arrangements in cases in which it would have been more advantageous for us to retain sole development and commercialization rights to such product candidate. If any of our product candidates receives marketing approval and we, or others, later discover that the drug is less effective than previously believed or causes undesirable side effects that were not previously identified, our ability to market the drug could be compromised. Clinical trials of our product candidates are conducted in carefully defined subsets of patients who have agreed to enter into clinical trials. Consequently, it is possible that our clinical trials may indicate an apparent positive effect of a product candidate that is greater than the actual positive effect, if any, or alternatively fail to identify undesirable side effects. If one or more of our product candidates receives regulatory approval, and we, or others, later discover that they are less effective than previously believed, or cause undesirable side effects, a number of potentially significant negative consequences could result, including: withdrawal, variation, suspension or limitation by regulatory authorities of approvals of such product; A 40 A product candidate is approved under 21 CFR 314 (Subpart H, accelerated approval) or we receive a conditional marketing authorization but required confirmatory trials may fail to verify clinical benefit or we may fail to fulfill requirements of the conditional marketing authorization; seizure of the product by regulatory authorities; recall of the product; restrictions on the marketing of the product or the manufacturing process for any component thereof; requirement by regulatory authorities of additional warnings on the label, such as a black box warning or contraindication; requirements that we implement a REMS, or comparable foreign strategies, or create a medication guide outlining the risks of such side effects for distribution to patients; commitment to expensive additional safety studies prior to approval or post-marketing studies required by regulatory authorities of such product; adverse impact on the product's competitiveness; initiation of regulatory investigations and government enforcement actions; initiation of legal action against us to hold us liable for harm caused to patients; and harm to our reputation and resulting harm to physician or patient acceptance of our products. Any of these events could prevent us from achieving or maintaining market acceptance of the particular product candidate, if approved, and could harm our business, financial condition, results of operations and prospects. We may become exposed to costly and damaging liability claims, either when testing our product candidates in the clinic or at the commercial stage, and our product liability insurance may not cover all damages from such claims. We are exposed to potential product liability and professional indemnity risks that are inherent in the research, development, manufacturing, marketing and use of pharmaceutical products. We currently have no products that have been approved for commercial sale. However, the current and future use of product candidates by us in clinical trials, and the sale of any approved products in the future, may expose us to liability claims. These claims might be made by patients who use the product, healthcare providers, pharmaceutical companies or others selling such products. Any claims against us, regardless of their merit, could be difficult and costly to defend or settle, and could compromise the market acceptance of our product candidates or any prospects for commercialization of our product candidates, if approved. For more information regarding the risks associated with intellectual property-related litigation, see "Risk Factors" "Risks Related to Our Intellectual Property." Although the clinical trial process is designed to identify and assess potential side effects, it is always possible that a drug, even after regulatory approval, may exhibit unforeseen or rare side effects. If any of our product candidates were to cause adverse side effects during clinical trials or after approval of the product candidate, we may be exposed to substantial liabilities. Physicians and patients may not comply with any warnings that identify known potential adverse effects and patients who should not use our product candidates. Although we maintain product liability insurance coverage, such insurance may not be adequate to cover all liabilities that we may incur. We may need to increase our insurance coverage each time we commence a clinical trial and if we successfully commercialize any product candidate. As the expense of insurance coverage is increasing, we may not be able to maintain insurance coverage at a reasonable cost or in an amount adequate to satisfy any liability that may arise. If a successful product liability claim or series of claims is brought against us for uninsured liabilities or in excess of insured liabilities, our assets may not be sufficient to cover such claims and our business operations could be impaired. Even if we receive Fast Track designation or granting of other FDA expedited programs, or other comparable foreign expedited programs, for any of our product candidates, there is no guarantee that such product candidates will experience a faster regulatory review or obtain regulatory approval. If a product is intended for the treatment of a serious or life-threatening condition and preclinical or clinical data demonstrate the potential to address an unmet medical need for this condition, the product sponsor may apply for Fast Track designation. The FDA has broad discretion whether or not to grant this designation, so even if we believe a particular product candidate is eligible for this designation, we cannot assure you that the FDA would decide to grant it. Even if we receive Fast Track designation for any of our product candidates, we may not experience a faster development process, review or approval compared to conventional FDA approval timelines, and the FDA may still decline to approve such product candidates. The FDA may rescind the Fast Track designation if it believes that the designation is no longer supported by data from our clinical development program or for any other reason. Similarly, A 41 A the FDA's other expedited drug development programs (e.g., Breakthrough Therapy, Accelerated Approval, Priority Review) do not guarantee a product candidate's faster regulatory review or regulatory approval. Although taltretectinib has been granted Breakthrough Therapy Designations by both the FDA and China's NMPA for the treatment of advanced or metastatic ROS1-positive NSCLC, such designations may be rescinded and may not lead to faster regulatory review or regulatory approval. The EMA has a similar program called PRIME. Even if we receive Orphan Drug designation for any of our product candidates, we may be unable to maintain the benefits associated with such designation, including the potential for market exclusivity. Regulatory authorities in some jurisdictions, including the United States and the EU, may designate drugs for relatively small patient populations as Orphan Drugs. Under the Orphan Drug Act, the FDA may designate a drug as an Orphan Drug if it is a drug intended to treat a rare disease or condition, which is generally defined as a patient population of fewer than 200,000 individuals in the United States, or a patient population greater than 200,000 in the United States where there is no reasonable expectation that the cost of developing the drug will be recovered from sales in the United States. In the United States, Orphan Drug designation entitles a party to financial incentives such as opportunities for grant funding towards clinical trial costs, tax credits for certain clinical trial costs and user-fee waivers. Generally, if a drug with an Orphan Drug designation subsequently receives the first marketing approval for the indication for which it has such designation, the drug is entitled to a period of marketing exclusivity, which precludes the FDA from approving another marketing application for the same drug and indication for that time period, except in limited circumstances. The applicable period is seven years in the United States. Even if we receive Orphan Drug designation for any of our product candidates, there is no guarantee that we will obtain approval or Orphan Drug exclusivity for such product candidates. Even if we obtain Orphan Drug exclusivity for any of our product candidates, that exclusivity may not effectively protect the product candidates from competition because different therapies can be approved for the same condition and the same therapy could be approved for different conditions. Even after an Orphan Drug is approved, the FDA can subsequently approve the same drug for the same condition if the FDA concludes that the later drug is clinically superior in that it is shown to be safer, more effective or makes a major contribution to patient care. Moreover, Orphan Drug exclusive marketing rights in the United States may be lost if the FDA later determines that the request for designation was materially defective or if the manufacturer is unable to assure sufficient quantity of the drug to meet the needs of patients with the rare disease or condition. Orphan Drug designation neither shortens the development time or regulatory review time of a drug nor gives the drug any advantage in the regulatory review or approval process. Risks Related to Commercialization of Our Product Candidates We have never commercialized a product candidate and we may lack the necessary expertise,

personnel and resources to successfully commercialize any of our products that receive regulatory approval on our own or together with collaborators. We have never commercialized a product candidate. Our operations to date have been limited to organizing and staffing our company, business planning, raising capital, undertaking preclinical studies of our product candidates and enrolling patients in clinical trials for our clinical stage product candidates. We currently have no sales force, marketing, manufacturing or distribution capabilities. To achieve commercial success of our product candidates, if any are approved, we will have to develop our own sales, marketing and manufacturing capabilities or outsource these activities to a third party. Factors that may affect our ability to commercialize our product candidates on our own include recruiting and retaining adequate numbers of effective sales and marketing personnel, persuading adequate numbers of physicians to prescribe our product candidates and other unforeseen costs associated with creating an independent sales and marketing organization. Developing a sales and marketing organization requires significant investment, is time-consuming and could delay the launch of our product candidates. We may not be able to build an effective sales and marketing organization in the U.S., the European Union or other key global markets. If we are unable to build our own distribution and marketing capabilities or to find suitable partners for the commercialization of our product candidates, we may have difficulties generating revenue from them. We face substantial competition, which may result in others discovering, developing or commercializing products before or more successfully than we do. The development and commercialization of new drug products is highly competitive. We face competition with respect to our current product candidates, and will face competition with respect to any product candidates that we may seek to develop or commercialize in the future, from major pharmaceutical, specialty pharmaceutical and biotechnology companies among others. We compete in the segments of the pharmaceutical, biotechnology and other related markets that develop immunotherapies for the treatment of cancer. There are other companies working to develop immunotherapies for the treatment of cancer including divisions of large pharmaceutical and biotechnology companies of various sizes. Some of these competitive products and therapies are based on scientific approaches that are the same as or similar to our approach, and others are based on entirely different approaches. Potential competitors also include academic institutions, government agencies and other public and private research organizations that conduct research, seek patent protection and establish collaborative arrangements for research, development, manufacturing and commercialization. ¶ 42 ¶ We are developing our initial product candidates for the treatment of cancer, and currently none of these therapies are approved. There are already a variety of available drug therapies marketed for cancer and some of the currently approved drug therapies are branded and subject to patent protection, and others are available on a generic basis. Many of these approved drugs are well established therapies and are widely accepted by physicians, patients and third-party payors. Insurers and other third-party payors may also encourage the use of generic products. We expect that if our product candidates are approved, they will be priced at a significant premium over competitive generic products. This may make it difficult for us to achieve our business strategy of replacing existing therapies with our product candidates. Our competitors may succeed in developing, acquiring or licensing, on an exclusive basis, products that are more effective or less costly than any product candidate that we may develop. In addition, most of these companies have substantially greater sales, marketing and other experience and reserves than we do. Competition may further increase as a result of advances in the commercial applicability of technologies for drug discovery and development and greater availability of capital for investment in cancer therapies. Established pharmaceutical companies may invest heavily to accelerate discovery and development of novel compounds or to in-license novel compounds that could make our product candidates less competitive. In addition, any new product that competes with an approved product must demonstrate compelling advantages in efficacy, convenience, tolerability and safety in order to overcome price competition and to be commercially successful. Accordingly, our competitors may succeed in obtaining patent protection, discovering, developing, receiving FDA approval for or commercializing drugs before we do, which would have an adverse impact on our business and results of operations. The availability of our competitors’ products could limit the demand and the price we are able to charge for any product candidate we commercialize, if any. The inability to compete with existing or subsequently introduced drugs would harm our business, financial condition, results of operations and prospects. Even if any of our product candidates receive marketing approval, they may fail to achieve the degree of market acceptance by physicians, patients, third-party payors and others in the medical community necessary for commercial success. If taltrectinib, fasusidenib, NUV-868, NUV-1511 and our other current and future product candidates receive marketing approval, whether as a single agent or in combination with other therapies, they may nonetheless fail to gain sufficient market acceptance by physicians, patients, third-party payors and others in the medical community. For example, current approved immunotherapies, and other cancer treatments like chemotherapy and radiation therapy, are well established in the medical community, and doctors may continue to rely on these therapies. If any of our product candidates do not achieve an adequate level of acceptance, we may not generate significant product revenues and we may never become profitable. The degree of market acceptance of our product candidates, if approved for commercial sale, will depend on a number of factors, including: ¶the ability of our product candidates to treat cancer, as compared with other available drugs, treatments or therapies; ¶the prevalence and severity of any adverse side effects associated with our product candidates; ¶limitations or warnings contained in the labeling approved for our product candidates by the FDA or comparable foreign regulatory authorities; ¶availability of alternative treatments; ¶the size of the target patient population, and the willingness of the target patient population to try new therapies and of physicians to prescribe these therapies; ¶the strength of marketing and distribution support and timing of market introduction of competitive products; ¶publicity for our product candidates and competing products and treatments; ¶pricing and cost effectiveness; ¶the effectiveness of our sales and marketing strategies; and ¶our ability to obtain sufficient third-party coverage and adequate reimbursement. The successful commercialization of our product candidates will depend in part on the extent to which governmental authorities and health insurers establish adequate coverage, reimbursement levels and pricing policies. Failure to obtain or maintain adequate coverage and reimbursement for our product candidates, if approved, could limit our ability to market those products and decrease our ability to generate revenue. The availability and adequacy of coverage and reimbursement by governmental healthcare programs such as Medicare and Medicaid, comparable foreign healthcare programs, private health insurers and other third-party payors are essential for most patients to be able to afford products such as our product candidates, if approved. Our ability to achieve acceptable levels of coverage and ¶ 43 ¶ reimbursement for products by governmental authorities, private health insurers and other organizations will have an effect on our ability to successfully commercialize our product candidates and, if desired, attract collaboration partners to invest in the development of our product candidates. Coverage under certain government programs, such as Medicare, Medicaid, the 340B drug pricing program and TRICARE, or comparable foreign healthcare programs, may not be available for certain of our product candidates. Assuming we obtain coverage for a given product by a third-party payor, the resulting reimbursement payment rates may not be adequate or may require co-payments that patients find unacceptably high. We cannot be sure that coverage and reimbursement in the U.S., the European Union or elsewhere will be available for any product that we may develop, and any reimbursement that may become available may be decreased or eliminated in the future. Third-party payors increasingly are challenging prices charged for pharmaceutical products and services, and many third-party payors may refuse to provide coverage and reimbursement for particular drugs when an equivalent generic drug, biosimilar or a less expensive therapy is available. It is possible that a third-party payor may consider our product candidates and other therapies as substitutable and only offer to reimburse patients for the less expensive product. Even if we show improved efficacy or improved convenience of administration with our product candidates, pricing of existing drugs may limit the amount we will be able to charge for our product candidates. These payors may deny or revoke the reimbursement status of a given product or establish prices for new or existing marketed products at levels that are too low to enable us to realize an appropriate return on our investment in product development. If reimbursement is not available or is available only at limited levels, we may not be able to successfully commercialize our product candidates, and may not be able to obtain a satisfactory financial return on products that we may develop. There is significant uncertainty related to the insurance coverage and reimbursement of newly approved products. In the U.S., third-party payors, including private and governmental payors, such as the Medicare and Medicaid programs, play an important role in determining the extent to which new drugs and biologics will be covered. The Medicare and Medicaid programs increasingly are used as models for how private payors and other governmental payors develop their coverage and reimbursement policies for drugs and biologics. Some third-party payors may require pre-approval of coverage for new or innovative devices or drug therapies before they will reimburse healthcare providers who use such therapies. It is difficult to predict at this time what third-party payors will decide with respect to the coverage and reimbursement for our product candidates. Obtaining and maintaining reimbursement status is time-consuming and costly. No uniform policy for coverage and reimbursement for products exists among third-party payors in the U.S. Therefore, coverage and reimbursement for products can differ significantly from payor to payor. As a result, the coverage determination process is often a time-consuming and costly process that will require us to provide scientific and clinical support for the use of our products to each payor separately, with no assurance that coverage and adequate reimbursement will be applied consistently or obtained in the first instance. Furthermore, rules and regulations regarding reimbursement change frequently, in some cases at short notice, and we believe that changes in these rules and regulations are likely. Moreover, increasing efforts by governmental and third-party payors in the U.S. and abroad to cap or reduce healthcare costs may cause such organizations to limit both coverage and the level of reimbursement for newly approved products and, as a result, they may not cover or provide adequate payment for our product candidates. We expect to experience pricing pressures in connection with the sale of any of our product candidates due to the trend toward managed healthcare, the increasing influence of health maintenance organizations, and additional legislative changes. The downward pressure on healthcare costs in general, particularly prescription drugs, has become very intense. As a result, increasingly high barriers are being erected to the entry of new products. The continuing efforts of the government, insurance companies, managed care organizations and other payors of health care services to contain or reduce costs of health care may adversely affect: ¶the demand for any products for which we may obtain regulatory approval; ¶our ability to set a price that we believe is fair for our products; ¶our ability to obtain coverage and reimbursement approval for a product; ¶our ability to generate revenues and achieve or maintain profitability; and ¶the level of taxes that we are required to pay. In addition, in case a drug product needs companion diagnostics, then companion diagnostic tests require coverage and reimbursement separate and apart from the coverage and reimbursement for the pharmaceutical or biological product. Similar challenges to obtaining coverage and reimbursement, applicable to pharmaceutical or biological products, will apply to companion diagnostics. Even if we obtain regulatory approval for our product candidates, they will remain subject to ongoing regulatory oversight. Even if we obtain regulatory approval for any of our product candidates, they will be subject to extensive and ongoing regulatory requirements for manufacturing processes, labeling, packaging, distribution, adverse event reporting, storage, advertising, promotion, sampling and record-keeping. These requirements include submissions of safety and other post-marketing information and reports, registration, as well as continued compliance with current Good Manufacturing Practice (“cGMP”), regulations and GCPs, for any clinical trials that we conduct post-approval, all of which may result in significant expense and limit our ability to commercialize ¶ 44 ¶ such products. In addition, any regulatory approvals that we receive for our product candidates may also be subject to limitations on the approved indicated uses for which the product may be marketed or to the conditions of approval, or contain requirements for potentially costly post-marketing testing, including Phase 4 clinical trials, and surveillance to monitor the safety and efficacy of the product candidate. In particular, a product may not be promoted for uses that are not approved by the FDA or such other regulatory authorities as reflected in the product’s approved labeling. If we receive marketing approval for any future product candidates we may develop, physicians may nevertheless prescribe it to their patients in a manner that is inconsistent with the approved label. However, if we are found to have promoted such off-label uses, we may become subject to significant liability. The FDA or comparable foreign regulatory authorities may also require a REMS, or comparable foreign regulatory strategies as a condition of approval of our product candidates, which could include requirements for a medication guide, physician communication plans or additional elements to ensure safe use, such as restricted distribution methods, patient registries and other risk minimization tools. The FDA’s and other regulatory authorities’ policies may change and additional government regulations may be enacted that could prevent, limit or delay regulatory approval of our product candidates. We cannot predict the likelihood, nature or extent of government regulation that may arise from future legislation or administrative action, either in the U.S. or abroad. If we are slow or unable to adapt to changes in existing requirements or the adoption of new requirements or policies, or if we are not able to maintain regulatory compliance, we may lose any marketing approval that we may have obtained and we may not achieve or sustain profitability. Moreover, if there are changes in the application of legislation or regulatory policies, or if problems are discovered with a product or our manufacture of a product, or if we or one of our distributors, licensees or co-marketers fails to comply with regulatory requirements, the regulators could take various actions. These include: ¶issuing warning or untitled letters; ¶seeking an injunction or imposing civil or criminal penalties or monetary fines; ¶suspension or imposition of restrictions on operations, including product manufacturing; ¶seizure or detention of products, refusal to permit the import or export of products or request that we initiate a product recall; ¶suspension, variation or withdrawal of our marketing authorizations; ¶suspension of any ongoing clinical trials; ¶refusal to approve pending applications or supplements to applications submitted by us; or ¶requiring us to conduct additional clinical trials, change our product labeling or submit additional applications for marketing authorization. If any of these events occurs, our ability to sell such product may be impaired, and we may incur substantial additional expense to comply with regulatory requirements, which could harm our business, financial condition, results of operations and prospects. If any of our product candidates are approved for marketing and commercialization and we are unable to establish sales and marketing capabilities or enter into agreements with third parties to sell and market our product candidates, we will be unable to successfully commercialize our product candidates if and when they are approved. We have no sales, marketing or distribution capabilities or experience. To achieve commercial success for any approved product for which we retain sales and marketing responsibilities, we must either develop a sales and marketing organization, which would be expensive and time consuming, or outsource these functions to other third parties. In the future, we may choose to build a focused sales and marketing infrastructure to sell, or participate in sales activities with our collaborators for, some of our product candidates if and when they are approved. There are risks involved with both establishing our own sales and marketing capabilities and entering into arrangements with third parties to perform these services. For example, recruiting and training a sales force is expensive and time consuming and could delay any product launch. If the commercial launch of a product candidate for which we recruit a sales force and establish marketing capabilities is delayed or does not occur for any reason, we would have prematurely or unnecessarily incurred these commercialization expenses. This may be costly, and our investment would be lost if we cannot retain or reposition our sales and marketing personnel. Factors that may inhibit our efforts to commercialize our product candidates on our own include: ¶our inability to recruit and retain adequate numbers of effective sales and marketing personnel; ¶the inability of sales personnel to obtain access to physicians or educate adequate numbers of physicians on the benefits of prescribing any future product candidates; ¶ 45 ¶ the lack of complementary products to be offered by sales personnel, which may put us at a competitive disadvantage relative to companies with more extensive product lines; and ¶unforeseen costs and expenses associated with creating an independent sales and marketing organization. If we enter into arrangements with third parties to perform sales, marketing and distribution services, our product revenues or the profitability of these product revenues to us are likely to be lower than if we were to market and sell any products that we develop ourselves. In addition, we may not be successful in entering into arrangements with third parties to sell and market our product candidates or may be unable to do so on terms that are favorable to us. In entering into third-party marketing or distribution arrangements, any revenue we receive will depend upon the efforts of the third parties and we cannot assure you that such third parties will establish adequate sales and distribution capabilities or devote the necessary resources and attention to sell and market our product candidates effectively. If we do not establish sales and marketing capabilities successfully, either on our own or in collaboration with third parties, we will not be successful in commercializing our product candidates. Risks Related to Our Dependence on Third Parties We rely on third parties to perform the chemistry work associated with our drug discovery and preclinical activities and to conduct our preclinical studies and future clinical trials, and our business could be substantially harmed if these third parties cease performing services or perform in an unsatisfactory manner. We do not have any laboratory facilities and have relied on CROs to perform most of the medicinal chemistry work associated with our drug discovery activities. We also do not currently have the ability to independently conduct preclinical studies or clinical trials without outside assistance. We have relied on CROs to conduct all of our preclinical studies to date and intends to conduct our future clinical trials by leveraging expertise and assistance from CROs as appropriate. We plan to rely upon medical institutions, clinical investigators, contract laboratories and other third parties, such as CROs, to conduct or assist us in conducting GCP-compliant clinical trials on our product candidates properly and on time, and may not currently have all of the necessary contractual relationships in place to do so. Once we have established contractual relationships with such third-party CROs, we will have only limited control over their actual performance of these activities. We and our CROs and other vendors are required to comply with cGMP, GCP, and good laboratory practice (GLP), which are regulations and guidelines enforced by the FDA, the competent authorities of EU Member States and any comparable foreign regulatory authorities for all of our product candidates in preclinical and clinical development. Regulatory authorities enforce

these regulations through periodic inspections of trial sponsors, principal investigators, clinical trial sites and other contractors. Although we will rely on CROs to conduct any current or planned GLP-compliant preclinical studies and GCP-compliant clinical trials and has limited influence over their actual performance, we remain responsible for ensuring that each of our preclinical studies and clinical trials is conducted in accordance with our investigational plan and protocol and applicable laws and regulations, and our reliance on the CROs does not relieve us of our regulatory responsibilities. If we or any of our CROs or vendors fail to comply with applicable regulations, the data generated in our preclinical studies and clinical trials may be deemed unreliable and the FDA, European Commission, MHRA, NMPA or any other comparable foreign regulatory authority may require us to perform additional preclinical studies and clinical trials before approving our marketing applications. We cannot assure you that upon inspection by a given regulatory authority, such regulatory authority will determine that all of our clinical trials comply with GCP regulations. In addition, our clinical trials must be conducted with products produced under cGMP requirements. We, our contract manufacturers, any future collaborators and their contract manufacturers could be subject to periodic unannounced inspections by the FDA, the competent authorities of EU Member States, the MHRA, the NMPA or other comparable foreign regulatory authorities, to monitor and ensure compliance with cGMP. Despite our efforts to audit and verify regulatory compliance, one or more of our third-party manufacturing vendors may be found on regulatory inspection by the FDA, the competent authorities of EU Member States, the MHRA, the NMPA or other comparable foreign regulatory authorities to be noncompliant with cGMP regulations. This may result in shutdown of the third-party vendor or invalidation of drug product lots or processes. In some cases, a product recall may be warranted or required, which would materially affect our ability to supply and market our drug products. Our failure to comply with these requirements may require us to repeat clinical trials, which would delay the regulatory approval process. While we or our CROs have or will have agreements governing their activities, we will not be able to control whether or not they devote sufficient time and resources to our future chemistry work and preclinical and clinical programs. These CROs may also have relationships with other commercial entities, including our competitors, for whom they may also be conducting clinical trials or other chemistry or drug discovery or development activities. We face the risk of potential unauthorized disclosure, infringement, misappropriation or other violation of our intellectual property by CROs, which may reduce our trade secret protection and allow our potential competitors, and other third parties, to access and exploit our proprietary technology. CROs also may use our proprietary information and intellectual property in such a way as to invite litigation or other intellectual property-related proceedings that could jeopardize or invalidate our proprietary information and intellectual property. If our CROs do not successfully carry out their contractual duties or obligations, fail to meet expected deadlines or if the quality or accuracy of the clinical data they obtain is compromised due to a 46 Å the failure to adhere to our clinical protocols or regulatory requirements or for any other reason, our clinical trials or other drug discovery or development activities may be extended, delayed or terminated, the clinical data generated in our clinical trials may be deemed unreliable, and we may not be able to obtain regulatory approval for, or successfully commercialize any product candidate that we develop. As a result, our financial results and the commercial prospects for any product candidate that we develop would be harmed, our costs could increase, and our ability to generate revenue could be delayed. If our relationships with our CROs were to terminate, we might not be able to enter into arrangements with alternative CROs or do so on commercially reasonable terms. Switching or adding additional CROs involves substantial cost and requires management time and focus, and could delay the discovery, development and commercialization of our product candidates. In addition, there is a natural transition period when a new CRO commences work. As a result, delays occur, which can negatively impact our ability to meet our desired clinical development timelines. Though we intend to carefully manage our relationships with our CROs, there can be no assurance that we will not encounter challenges or delays in the future or that these delays or challenges will not have a negative impact on our business and financial condition. We do not have our own manufacturing capabilities and will rely on third parties to produce clinical and commercial supplies of talectrectinib, safusidenib, NUV-868, NUV-1511 and our other current and future product candidates. We have limited experience in drug formulation and manufacturing and do not own or operate, and we do not expect to own or operate, facilities for drug manufacturing, storage, distribution or testing. To date, we have obtained active pharmaceutical ingredients (ÅPisÅ) and drug product for our investigational products mostly from single-source third-party CMOs. We are in the process of developing our supply chain for each of our investigational products and intend to put in place framework agreements under which CMOs will generally provide us with necessary quantities of API and drug product on a project-by-project basis based on our development needs. We seek to use a different CMO for each investigational product and will consider further diversification of drug product and supply organizations as circumstances warrant. Third-party CMOs may be unable or unwilling to supply us with sufficient clinical and commercial grade quantities of our clinical materials due to production shortages or other supply interruptions resulting from health epidemics, because they are purchased by one of our competitors or another company that decides not to continue supplying us with these materials, or for other reasons. If one or more of these events occur and we are unable to timely establish an alternate supply from one or more third-party CMOs, we could experience delays in our development efforts as we locate and qualify new manufacturers. Under such circumstances, we may be required to receive drug substance for use on a purchase order basis, and as such, there can be no assurance that we actually receive sufficient quantities. See also the risk factor titled ÅOur business, operations and clinical development plans and timelines and supply chain could be adversely affected by the effects of health epidemics, on the manufacturing, clinical trial and other business activities performed by us or by third parties with whom we conduct business, including our CMOs, CROs, shippers and others.Å Further, our reliance on third-party manufacturers exposes us to risks beyond our control, including the risk of: Åinability to meet our product specifications and quality requirements consistently; Ådelay or inability to procure or expand sufficient manufacturing capacity; Åmanufacturing and quality issues, including related to scale-up of manufacturing; Åcosts and validation of new equipment and facilities required for additional scale-up; Åfailure of the manufacturer to comply with cGMP and similar foreign standards; Åinability to negotiate manufacturing agreements with third parties on commercially reasonable terms; Återmination or nonrenewal of manufacturing agreements with third parties in a manner or at a time that is costly or damaging to us; Åreliance on a limited number of sources, and in some cases, single sources for components, such that if we are unable to secure a sufficient supply of these drug components, we will be unable to manufacture and sell our product candidates in a timely fashion, in sufficient quantities or under acceptable terms; Ålack of qualified backup suppliers for those components that are currently purchased from a sole or single source supplier; Åoperations of our third-party manufacturers or suppliers could be disrupted by conditions unrelated to our business or operations, including the bankruptcy of the manufacturer or supplier or the issuance of a FDA Form 483 notice, warning letter, or cease and desist order; Åcarrier disruptions or increased costs that are beyond our control; and Åfailure to deliver our products under specified storage conditions and in a timely manner. Å 47 Å Some of these events could be the basis for FDA or comparable foreign regulatory authority action, including injunction, recall, seizure or total or partial suspension of production. In addition, our third-party manufacturers and suppliers are subject to FDA inspection and may be subject to inspections from comparable foreign regulatory authorities from time to time. Failure by our third-party manufacturers and suppliers to pass such inspections and otherwise satisfactorily complete the FDA approval regimen, or comparable foreign regulatory authorities' approval regimen, with respect to our product candidate may result in regulatory actions such as the issuance of FDA Form 483 notices of observations, warning letters or injunctions or the loss of operating licenses. In addition, our third-party manufacturers and suppliers are subject to numerous environmental, health and safety laws and regulations, including those governing the handling, use, storage, treatment and disposal of waste products, and failure to comply with such laws and regulations could result in significant costs associated with civil or criminal fines and penalties for such third parties. Based on the severity of the regulatory action, our clinical or commercial supply of drug and packaging and other services could be interrupted or limited, which could harm our business. In addition, our CMOs are or may be engaged with other companies to supply and manufacture materials or products for such companies, which also exposes our suppliers and manufacturers to regulatory risks for the production of such materials and products. As a result, failure to meet the regulatory requirements for the production of those materials and products may also affect the regulatory clearance of a contract supplier's or manufacturer's facility. If the FDA or a comparable foreign regulatory authority finds these facilities unsatisfactory in compliance with applicable regulations, does not approve these facilities for the supply or manufacture of our product candidates, or if it withdraws its approval in the future, we may need to find alternative supply or manufacturing facilities, which would negatively impact our ability to develop, obtain regulatory approval of our market our product candidates, if approved. As we prepare for later-stage clinical trials and potential commercialization, we will need to take steps to increase the scale of production of our product candidates, which may include transferring production to new third-party suppliers or manufacturers. In order to conduct larger or late-stage clinical trials for our product candidates and supply sufficient commercial quantities of the resulting drug product and our components, if that product candidate is approved for sale, our CMOs and suppliers will need to produce our product candidates in larger quantities, more cost effectively and, in certain cases, at higher yields than they currently achieve. These third-party contractors may not be able to successfully increase the manufacturing capacity for any such product candidates in a timely or cost-effective manner or at all. Significant scale up of manufacturing may require additional processes, technologies and validation studies, which are costly, may not be successful and which the FDA and comparable foreign regulatory authorities must review and approve. In addition, quality issues may arise during those scale-up activities because of the inherent properties of a product candidate itself or of a product candidate in combination with other components added during the manufacturing and packaging process, or during shipping and storage of the APIs or the finished product. If our third-party CMOs are unable to successfully scale up the manufacture of any of our product candidates in sufficient quality and quantity and at commercially reasonable prices, and we are unable to find one or more replacement suppliers or manufacturers capable of production at a substantially equivalent cost in substantially equivalent volumes and quality, and we are unable to successfully transfer the processes on a timely basis, the development of that product candidate and regulatory approval or commercial launch for any resulting products may be delayed, or there may be a shortage in supply, either of which could significantly harm our business, financial condition, results of operations and prospects. Failure to maintain the License Agreement between Daiichi Sankyo Company, Limited and AnHeart Therapeutics Inc., dated December 7, 2018, as amended (the ÅTalectrectinib In-License AgreementÅ) and the License Agreement between Daiichi Sankyo Company, Limited and AnHeart Therapeutics Inc., dated September 7, 2020 (the ÅSafusidenib In-License AgreementÅ) could negatively impact our business. Pursuant to the terms of the Talectrectinib In-License Agreement and the Safusidenib In-License Agreement, we received certain exclusive licenses to develop, manufacture and commercialize talectrectinib and safusidenib, respectively. Consequently, our ability to develop and commercialize talectrectinib and safusidenib depends on our ability to maintain these agreements with Daiichi Sankyo. We are subject to a number of other risks associated with our dependence on the Talectrectinib In-License Agreement and the Safusidenib In-License Agreement, including: ÅOur obligations to make certain milestone and royalty payments; ÅOur obligation to use commercially reasonable efforts to perform certain development and commercialization activities and to achieve certain milestones; ÅCertain obligations not to develop or commercialize products that compete with talectrectinib or safusidenib; and ÅPotential disputes between us and Daiichi Sankyo, including disagreements regarding the Talectrectinib In-License Agreement and the Safusidenib In-License Agreement. If either the Talectrectinib In-License Agreement or the Safusidenib In-License Agreement is terminated early, we may be unable to pursue continued development, manufacture and commercialization of talectrectinib or safusidenib. Å 48 Å If we are not able to establish and maintain collaborations, we may have to alter some of our future development and commercialization plans. Our product development programs and the potential commercialization of our product candidates will require substantial additional capital to fund expenses. The commercial rights to talectrectinib have been out-licensed in China and Japan, and we may enter into other collaboration agreements with pharmaceutical and biotechnology companies for the future development and potential commercialization of our product candidates. We will likely have limited control over the amount and timing of resources that our current and future collaborators dedicate to the development or commercialization of any product candidates we may seek to develop and commercialize with them. We cannot predict the success of any current or future collaboration that we may enter into. We face significant competition in seeking appropriate collaborators, and a number of more established companies may also be pursuing strategies to license or acquire third-party intellectual property rights that we may consider attractive. These established companies may have a competitive advantage over us due to their size, financial resources and greater clinical development and commercialization experience and capabilities. In addition, companies that perceive us to be a competitor may be unwilling to assign or license rights to us. Whether we reach a definitive agreement for a collaboration will depend, among other things, upon our assessment of the collaborator's resources and expertise, the terms and conditions of the proposed collaboration and the proposed collaborator's evaluation of a number of factors. Those factors may include the design or results of clinical trials, the likelihood of approval by the FDA, European Commission, MHRA, NMPA or other similar foreign regulatory authorities, the potential market for the subject product candidate, the costs and complexities of manufacturing and delivering such product candidate to patients, competing products, the existence of uncertainty with respect to our ownership of technology, which can exist if there is a challenge to such ownership without regard to the merits of the challenge, and industry and market conditions generally. The collaborator may also consider alternative product candidates or technologies for similar indications that may be available to collaborate on and whether such a collaboration could be more attractive than the one with us for our product candidate. We may also be restricted under future license agreements from entering into agreements on certain terms with potential collaborators. Collaborations are complex and time-consuming to negotiate and document. In addition, there have been a significant number of recent business combinations among large pharmaceutical companies that have resulted in a reduced number of potential future collaborators. We may not be able to negotiate collaboration agreements on a timely basis, on acceptable terms, or at all. Even if we are able to obtain a license to intellectual property of interest, we may not be able to secure exclusive rights, in which case others could use the same rights and compete with us. Our collaboration partners, if any, may not prioritize our product candidates or otherwise not effectively pursue the development of our product candidates which may delay, reduce or terminate the development of such product candidate, reduce or delay its development program or delay its potential commercialization. Further if we are unable to successfully obtain rights to required third-party intellectual property rights or maintain and protect the existing intellectual property rights we have, we may have to delay, reduce or terminate the development of our product candidates, reduce or delay its development program or one or more of our other development programs, delay its potential commercialization or reduce the scope of any sales or marketing activities or increase our expenditures and undertake development or commercialization activities at our own expense. Doing so will likely harm our ability to execute our business plans. If we elect to increase our expenditures to fund development or commercialization activities on our own, we may need to obtain additional capital, which may not be available to us on acceptable terms or at all. If we do not have sufficient funds, we may not be able to further develop our product candidates or bring them to market and generate product revenue. Risks Related to Regulatory Compliance Enacted and future legislation may increase the difficulty and cost for us to obtain marketing approval of and commercialize our product candidates and affect the prices we may charge for such product candidates. The U.S. and many foreign jurisdictions have enacted or proposed legislative and regulatory changes affecting the healthcare system that could prevent or delay marketing approval of our product candidates, restrict or regulate post-approval activities and affect our ability to profitably sell any product for which we obtain marketing approval. The Patient Protection and Affordable Care Act of 2010, as amended by the Health Care and Education Reconciliation Act of 2010 (collectively, the ÅAffordable Care ActÅ), includes measures that have significantly changed the way healthcare is financed by both governmental and private insurers. There have been judicial, executive and congressional challenges to certain aspects of the Affordable Care Act. In 2021 the U.S. Supreme Court dismissed a challenge on procedural grounds that argued the Affordable Care Act is unconstitutional in its entirety because the Åindividual mandateÅ was repealed by Congress. In addition, there have been a number of health reform initiatives by the Biden administration that have impacted the Affordable Care Act. For example, the Inflation Reduction Act of 2022 (ÅIRAÅ), extends enhanced subsidies for individuals purchasing health insurance coverage in Affordable Care Act marketplaces through plan year 2025. The IRA also eliminates the Ådonut holeÅ under the Medicare Part D program beginning in 2025 by significantly lowering the beneficiary maximum out-of-pocket cost and through a newly established manufacturer discount program. It is possible that the Affordable Care Act will be subject to judicial or Congressional challenges in the future. It is unclear how any such challenges and the healthcare reform measures of the Biden administration will impact the Affordable Care Act and our business. We continue to evaluate the effect that the Affordable Care Act and its possible repeal and replacement has on our business. Å 49 Å In addition, other legislative changes have been proposed and adopted since the Affordable Care Act was enacted. For example, the Budget Control Act of 2011, among other things, created the Joint Select Committee on Deficit Reduction to recommend to Congress proposals in spending reductions. The Joint Select Committee on Deficit Reduction did not achieve a targeted deficit reduction, which triggered the legislation's automatic reduction to several government programs. This includes aggregate reductions to Medicare payments to providers of, on average, 2% per fiscal year until 2032 unless Congress takes additional action. Additionally, the American Rescue Plan Act of 2021 eliminates the statutory Medicaid drug rebate cap, previously set at 100% of a drug's average manufacturer price, for single source and innovator multiple source drugs, effective January 1, 2024.

Recently, there has been increasing legislative and enforcement interest in the U.S. with respect to specialty drug pricing practices. Specifically, there have been several recent U.S. presidential executive orders, congressional inquiries and legislation designed to, among other things, bring more transparency to drug pricing, reduce the cost of prescription drugs under Medicare, review the relationship between pricing and manufacturer patient programs and reform government program reimbursement methodologies for drugs. For example, at the federal level, the Trump administration used several means to propose or implement drug pricing reform, including through federal budget proposals, executive orders and policy initiatives. In 2021, the Biden administration released an executive order, “Promoting Competition in the American Economy,” with multiple provisions aimed at prescription drugs. In response to Biden’s executive order, later in 2021, the Department of Health and Human Services (HHS) released a Comprehensive Plan for Addressing High Drug Prices that outlines principles for drug pricing reform and sets out a variety of potential legislative policies that Congress could pursue as well as potential administrative actions HHS can take to advance these principles. In addition, the IRA, among other things, (1) directs HHS to negotiate the price of certain single-source drugs and biologics covered under Medicare and (2) imposes rebates under Medicare Part B and Medicare Part D to penalize price increases that outpace inflation. These provisions take effect progressively starting in fiscal year 2023. In 2023, HHS announced the list of the first ten drugs that will be subject to price negotiations, although the Medicare drug price negotiation program is currently subject to legal challenges. It is currently unclear how the IRA will be implemented but is likely to have a significant impact on the pharmaceutical industry. In response to the Biden administration’s 2022 executive order, in 2023, HHS released a report outlining three new models for testing by the Center for Medicare and Medicaid Innovation which will be evaluated on their ability to lower the cost of drugs, promote accessibility, and improve quality of care. It is unclear whether the models will be utilized in any health reform measures in the future. Further, in 2023, the Biden administration announced an initiative to control the price of prescription drugs through the use of march-in rights under the Bayh-Dole Act. Also in 2023, the National Institute of Standards and Technology published for comment a Draft Interagency Guidance Framework for Considering the Exercise of March-In Rights which for the first time includes the price of a product as one factor an agency can use when deciding to exercise march-in rights. While march-in rights have not previously been exercised, it is uncertain if that will continue under the new framework. At the state level, legislatures have increasingly passed legislation and implemented regulations designed to control pharmaceutical and biological product pricing, including price or patient reimbursement constraints, discounts, restrictions on certain product access and marketing cost disclosure and transparency measures, and, in some cases, designed to encourage importation from other countries and bulk purchasing. For example, in January 2024, the FDA approved Florida’s Section 804 Importation Program (SIP) proposal to import certain drugs from Canada for specific state healthcare programs. It is unclear how this program will be implemented, including which drugs will be chosen, and whether it will be subject to legal challenges in the United States or Canada. Other states have also submitted SIP proposals that are pending review by the FDA. Any such approved importation plans, when implemented, may result in lower drug prices for products covered by those programs. We are unable to predict the future course of federal or state healthcare legislation in the U.S. directed at broadening the availability of healthcare and containing or lowering the cost of healthcare. These and any further changes in the law or regulatory framework that reduce our revenue or increase our costs could also have a material and adverse effect on our business, financial condition and results of operations. We expect that the healthcare reform measures that have been adopted and may be adopted in the future, may result in more rigorous coverage criteria and in additional downward pressure on the price that we receive for any approved product and could seriously harm our future revenues. Any reduction in reimbursement from Medicare or other government programs may result in a similar reduction in payments from private payors. The implementation of cost containment measures or other healthcare reforms may prevent us from being able to generate revenue, attain profitability or commercialize our products. Our business operations and current and future relationships with investigators, healthcare professionals, consultants, third-party payors and customers will be subject, directly or indirectly, to federal and state healthcare fraud and abuse laws, false claims laws, transparency laws, health information privacy and security laws and other healthcare laws and regulations, including comparable foreign healthcare laws and regulations. If we are unable to comply, or have not fully complied, with such laws, we could face substantial penalties. Although we do not currently have any products on the market, our current and future operations may be, directly or indirectly through our prescribers, customers and third-party payors, subject to various U.S. federal and state healthcare laws and regulations. Healthcare providers and others play a primary role in the recommendation and prescription of any products for which we obtain a 50 A marketing approval. These laws may impact, among other things, our current business operations, including our clinical research activities, and proposed sales, marketing and education programs and constrain the business of financial arrangements and relationships with healthcare providers and other parties through which we may market, sell and distribute our products for which we obtain marketing approval. The laws that may affect our ability to operate include: the U.S. federal Anti-Kickback Statute, which prohibits, among other things, persons or entities from knowingly and willfully soliciting, offering, receiving or paying any remuneration (including any kickback, bribe or certain rebates), directly or indirectly, overtly or covertly, in cash or in kind, to induce or reward either the referral of an individual for, or the purchase, lease, order or recommendation of, any good, facility, item or service, for which payment may be made, in whole or in part, under U.S. federal and state healthcare programs such as Medicare and Medicaid. A person or entity does not need to have actual knowledge of the statute or specific intent to violate it in order to have committed a violation; the U.S. federal false claims, including the False Claims Act, which can be enforced through whistleblower actions, and Civil Monetary Penalties Laws, which, among other things, impose criminal and civil penalties against individuals or entities for knowingly presenting, or causing to be presented, to the U.S. federal government, claims for payment or approval that are false or fraudulent, knowingly making, using or causing to be made or used, a false record or statement material to a false or fraudulent claim, or from knowingly making a false statement to avoid, decrease or conceal an obligation to pay money to the U.S. federal government. In addition, the government may assert that a claim including items and services resulting from a violation of the U.S. federal Anti-Kickback Statute constitutes a false or fraudulent claim for purposes of the False Claims Act; the U.S. federal Health Insurance Portability and Accountability Act of 1996, (HIPAA), which imposes criminal and civil liability for, among other things, knowingly and willfully executing, or attempting to execute, a scheme to defraud any healthcare benefit program, or knowingly and willfully falsifying, concealing or covering up a material fact or making any materially false statement, in connection with the delivery of, or payment for, healthcare benefits, items or services; similar to the U.S. federal Anti-Kickback Statute, a person or entity does not need to have actual knowledge of the statute or specific intent to violate it in order to have committed a violation; HIPAA, as amended by the Health Information Technology for Economic and Clinical Health Act of 2009 (HITECH), and its implementing regulations, which imposes certain obligations, including mandatory contractual terms, with respect to safeguarding the privacy, security and transmission of individually identifiable health information without appropriate authorization by covered entities, i.e. health plans, healthcare clearinghouses and certain healthcare providers, as well as their business associates and covered subcontractors that perform certain services for or on their behalf involving the use or disclosure of individually identifiable health information; the U.S. Federal Food, Drug and Cosmetic Act, which prohibits, among other things, the adulteration or misbranding of drugs, biologics and medical devices; the U.S. federal legislation commonly referred to as Physician Payments Sunshine Act, enacted as part of the Affordable Care Act, and its implementing regulations, which requires certain manufacturers of drugs, devices, biologics and medical supplies that are reimbursable under Medicare, Medicaid or the Children’s Health Insurance Program to report annually to the Centers for Medicare & Medicaid Services (“CMS”) information related to certain payments and other transfers of value to physicians (defined to include doctors, dentists, optometrists, podiatrists and chiropractors), certain other healthcare professionals (such as physician assistants and nurse practitioners), and teaching hospitals, as well as ownership and investment interests held by the physicians described above and their immediate family members; analogous state laws and regulations, including: state anti-kickback and false claims laws, which may apply to our business practices, including, but not limited to, research, distribution, sales and marketing arrangements and claims involving healthcare items or services reimbursed by any third-party payor, including private insurers; state laws that require pharmaceutical companies to comply with the pharmaceutical industry’s voluntary compliance guidelines and the relevant compliance guidance promulgated by the U.S. federal government, or otherwise restrict payments that may be made to healthcare providers and other potential referral sources; state laws and regulations that require drug manufacturers to file reports relating to pricing and marketing information, which requires tracking gifts and other remuneration and items of value provided to healthcare professionals and entities; state and local laws requiring the registration of pharmaceutical sales representatives; and state laws governing the privacy and security of health information in certain circumstances, many of which differ from each other in significant ways and often are not preempted by HIPAA, thus complicating compliance efforts; and European and other foreign law equivalents of each of the laws, including reporting requirements detailing interactions with and payments to healthcare providers. Ensuring that our internal operations and future business arrangements with third parties comply with applicable healthcare laws and regulations will involve substantial costs. It is possible that governmental authorities will conclude that our business practices do not comply with current or future statutes, regulations, agency guidance or case law involving applicable fraud and abuse or other healthcare laws and regulations. If our operations are found to be in violation of any of the laws described above or any other governmental laws and regulations that may apply to us, we may be subject to significant penalties, including civil, criminal and administrative penalties, damages, fines, exclusion from U.S. government funded healthcare programs, such as Medicare and Medicaid, or similar programs in other countries or jurisdictions, disgorgement, imprisonment, contractual damages, reputational harm, diminished profits, additional reporting requirements and oversight if we become subject to a corporate integrity agreement or similar agreement to resolve allegations of non-compliance with these laws and the delay, reduction, termination or restructuring of our operations. Further, defending against any such actions can be costly and time-consuming, and may require significant financial and personnel resources. Therefore, even if we are successful in defending against any such actions that may be brought against us, our business may be impaired. If any of the physicians or other providers or entities with whom we expect to do business is found to not be in compliance with applicable laws, they may be subject to significant criminal, civil or administrative sanctions, including exclusions from government funded healthcare programs and imprisonment. If any of the above occur, it could adversely affect our ability to operate our business and our results of operations. We and the third parties with whom we work are subject to stringent and evolving U.S. and foreign laws, regulations, and rules, contractual obligations, industry standards, policies and other obligations related to data privacy and security. Our (or the third parties with whom we work) actual or perceived failure to comply with such obligations could lead to regulatory investigations or actions; litigation (including class claims) and mass arbitration demands; fines and penalties; disruptions of our business operations; reputational harm; loss of revenue or profits; loss of revenue or profits; and other adverse business consequences. In the ordinary course of business, we collect, receive, store, process, generate, use, transfer, disclose, make accessible, protect, secure, dispose of, transmit, and share (collectively, process) personal data and other sensitive information, including proprietary and confidential business information, trade secrets, intellectual property, information we collect about trial participants in connection with clinical trials, and sensitive third-party information (collectively, sensitive data). Our processing activities subject us to numerous data privacy and security obligations, such as various laws, regulations, guidance, industry standards, external and internal privacy and security policies, contractual requirements and other obligations relating to data privacy and security. In the United States, federal, state, and local governments have enacted numerous data privacy and security laws, including data breach notification laws, personal data privacy laws, consumer protection laws (e.g., Section 5 of the Federal Trade Commission Act), and other similar laws (e.g., wiretapping laws). For example, HIPAA, as amended by HITECH, imposes specific requirements relating to the privacy, security, and transmission of individually identifiable protected health information. Additionally, in the past few years, numerous U.S. states including California, Virginia, Colorado, Connecticut, and Utah have enacted comprehensive privacy laws that impose certain obligations on covered businesses, including providing specific disclosures in privacy notices and affording residents with certain rights concerning their personal data. These state laws allow for statutory fines for noncompliance. For example, the California Consumer Privacy Act of 2018, or CCPA, applies to personal data of California residents and requires businesses subject to the CCPA to provide specific disclosures in privacy notices and respond to requests of such individuals to exercise certain privacy rights. Although there are minimum revenue thresholds for entities to be subject to these laws and there are limited exemptions for clinical trial data under the CCPA and similar US state comprehensive privacy laws, such laws may impact (possibly significantly) our business activities depending on how they are interpreted, should we become subject to the CCPA or other such state comprehensive privacy laws in the future. Similar laws are being considered in other states, as well as at the federal and local levels, and we expect more laws related to personal data to become effective in the future. These developments may further complicate compliance efforts and increase our legal risk and compliance costs. Outside the United States, an increasing number of laws, regulations, and industry standards may govern data privacy and security. For example, the European Union’s General Data Protection Regulation, or EU GDPR, the United Kingdom’s GDPR, or UK GDPR, Australia’s Privacy Act, and China’s Personal Information Protection Law, or PIPL, impose strict requirements for processing personal data. For example, under GDPR, companies may face private litigation related to processing of personal data brought by classes of data subjects or consumer protection organizations authorized at law to represent their interests, temporary or definitive prohibitions on data processing and other corrective actions, or fines of up to the greater of 20 million Euros under the EU GDPR / 17.5 million pounds under the UK GDPR, or 4% of their worldwide annual revenue, whichever is higher. In addition, we may be unable to transfer personal data from Europe and other jurisdictions to the United States or other countries due to data localization requirements or limitations on cross-border data flows. Europe and other jurisdictions have enacted laws requiring data to be localized or limiting the transfer of personal data to other countries. In particular, the European Economic Area, or EEA, and the UK have significantly restricted the transfer of personal data to the United States and other countries whose privacy laws it generally believes are inadequate. Other jurisdictions may adopt similarly stringent interpretations of their data localization and cross-border data transfer laws. Although there are currently various mechanisms that may be used to transfer personal data from the EEA and UK to the United States in compliance with law, such as the EEA’s standard contractual clauses, the UK’s International Data Transfer Agreement / Addendum, and the EU-U.S. Data Privacy Framework and the UK extension thereto (which allows for transfers to relevant U.S.-based organizations who self-certify compliance and participate in the Framework), these mechanisms are subject to legal challenges, and there is no assurance that we can satisfy or rely on these measures to lawfully transfer personal data to the United States. If there is no lawful manner for us to transfer personal data from the EEA, the UK, or other jurisdictions to the United States, or if the requirements for a legally-compliant transfer are too onerous, we could face significant adverse consequences, including the interruption or degradation of our operations, the need to relocate part or all of our business or data processing activities to other jurisdictions (such as Europe) at significant expense, increased exposure to regulatory actions, substantial fines and penalties, the inability to transfer data and work with partners, vendors and other third parties, and injunctions against our processing or transferring of personal data necessary to operate our business. Additionally, companies that transfer personal data out of the EEA and UK to other jurisdictions, particularly to the United States, are subject to increased scrutiny from regulators, individual litigants, and activist groups. Some European regulators have ordered certain companies to suspend or permanently cease certain transfers of personal data out of Europe for allegedly violating the EU GDPR’s cross-border data transfer limitations. We may also publish privacy policies, marketing materials, and other statements, such as compliance with certain certifications or self-regulatory principles, regarding data privacy and security. If these policies, materials, or statements are found to be deficient, lacking in transparency, deceptive, unfair, or misrepresentative of our practices, we may be subject to investigation, enforcement actions by regulators, or other adverse consequences. Obligations related to data privacy and security (and individuals’ data privacy expectations) are quickly changing, becoming increasingly stringent, and creating uncertainty. Additionally, these obligations may be subject to differing applications and interpretations, which may be inconsistent or conflict among jurisdictions. Preparing for and complying with these obligations requires us to devote significant resources and may necessitate changes to our services, information technologies, systems, and practices and to those of any third parties that process personal data on our behalf. We may at times fail (or be perceived to have failed) in our efforts to comply with our data privacy and security obligations. Moreover, despite our efforts, our personnel or third parties with whom we work (such as contract research organizations and clinical trial sites) may fail (or be perceived to have failed) to comply with such obligations, which could negatively impact our business operations. If we or the third parties with whom we work fail, or are perceived to have failed, to address or comply with applicable data privacy and security obligations, we could face significant consequences, including but not limited to, government enforcement actions (e.g., investigations, fines, penalties, audits, inspections, and similar); litigation (including class-action claims) and mass arbitration demands; additional reporting requirements and/or oversight; bans or restrictions on processing personal data (including clinical trial data); orders to destroy or not use personal data; and imprisonment of company officials. In particular, plaintiffs have become increasingly more active in bringing privacy-related claims against companies, including class claims and mass arbitration demands. Some of these claims allow for the recovery of statutory damages on a per violation basis, and, if viable, carry the potential for monumental statutory damages,

depending on the volume of data and the number of violations. Any of these events could have a material adverse effect on our reputation, business, or financial condition, including but not limited to: interruptions or stoppages in our business operations (including, as relevant, clinical trials); inability to process sensitive data or to operate in certain jurisdictions; limited ability to develop or commercialize our products; expenditure of time and resources to defend any claim or inquiry; adverse publicity; or substantial changes to our business model or operations. Risks Related to Our Intellectual Property If we are unable to obtain, maintain, protect and enforce sufficient patent and other intellectual property rights for our product candidates and technology, or if the scope of patent and other intellectual property rights obtained is not sufficiently broad, we may not be able to compete effectively in our market. Our success depends in significant part on our ability and the ability of any licensors and collaborators to obtain, maintain, protect and enforce patents and other intellectual property rights with respect to our product candidates and technology and to operate our business without infringing, misappropriating or otherwise violating the intellectual property rights of others. The patent prosecution process is uncertain, expensive and time-consuming. We and our current or future licensors, licensees or collaborators may not be able to prepare, file and prosecute all necessary or desirable patent applications at a reasonable cost or in a timely manner. It is also possible that we or our future licensors will fail to identify patentable aspects of our research and development output in time to obtain patent protection or fail to file patent applications covering inventions made in the course of development and commercialization activities before a competitor or another third party files a patent application covering, or publishes information disclosing, a same or similar, independently-developed invention. Such competitor's or other third party's patent application or published information may pose obstacles to or prohibit our ability to obtain patent protection or limit the scope of the patent protection we may obtain. Although we enter into non-disclosure and confidentiality agreements with parties who have access to confidential or patentable aspects of our research and development output, such as our employees, collaborators, CROs, contract manufacturers, consultants, advisors and other third parties, any of these parties may breach the agreements and disclose such output before a patent application is filed, thereby jeopardizing our ability to seek patent protection in certain jurisdictions. In addition, publications of discoveries in the scientific literature often lag behind actual discoveries, and patent applications in the U.S. and other jurisdictions are 53 Å typically not published until approximately 18 months after filing, or in some cases not at all. Therefore, we cannot be certain that we or our future licensors were the first to conceive the inventions claimed in our owned or licensed patents or pending patent applications, or were the first to file for patent protection of such inventions. The patent position of biotechnology and pharmaceutical companies generally is uncertain, involves complex legal and factual questions and is the subject of much litigation. As a result, the issuance, scope, validity, enforceability and commercial value of our and our current or future licensors' patent rights are uncertain. Our and our licensors' pending and future patent applications may not mature into patents or result in issued patents that protect our technology or product candidates, in whole or in part, or effectively exclude others from commercializing competitive technologies and product candidates. The patent examination process may require us or our licensors to narrow the scope of the claims of our pending and future patent applications, and therefore, even if such patent applications issue as patents, they may not issue in a form that will provide us with any meaningful protection, prevent competitors or other third parties from competing with us or otherwise provide us with any competitive advantage. Our and our licensors' patent applications cannot be enforced against third parties practicing the technology claimed in such applications unless and until a patent issues from such applications, and then only to the extent the issued claims cover such technology. Any patents that we hold or in-license may be challenged or, circumvented by third parties or narrowed, invalidated or held unenforceable in litigation or post-grant proceedings. Consequently, we do not know whether any of our product candidates will be protectable or remain protected by valid and enforceable patents. Our competitors or other third parties may be able to circumvent our patents by developing similar or alternative technologies or products in a non-infringing manner. Any of the foregoing could harm our competitive position, business, financial condition, results of operations and prospects. The patent protection we obtain for our product candidates and technology may be challenged or not sufficient to provide us with any competitive advantage. Even if our owned or licensed patent applications issue as patents, the issuance of any such patents is not conclusive as to their inventorship, scope, validity or enforceability, and such patents may be challenged, invalidated, narrowed or held to be unenforceable, including in the courts or patent offices in the U.S. and abroad, or circumvented. We may be subject to a third-party preissuance submission of prior art to the U.S. Patent and Trademark Office (the "USPTO"), a federal court or equivalent foreign bodies, or become involved in opposition, derivation, revocation, re-examination, post-grant and inter partes review or interference proceedings, or other similar proceedings, challenging our patent rights or the patent rights of others. An adverse determination as a result of any such submission, proceeding or litigation could reduce the scope of, invalidate, or render unenforceable, our patent rights, allow third parties to commercialize our technology or products and compete directly with us, without payment to us, or result in our inability to manufacture or commercialize products without infringing third-party patent rights. Moreover, we, or one of our licensors, may have to participate in interference or derivation proceedings declared by the USPTO to determine priority or ownership of invention or in post-grant challenge proceedings, such as oppositions in a foreign patent office, that challenge priority of invention or other features of patentability. Such proceedings and any other patent challenges may result in loss of patent rights, loss of exclusivity, loss of priority or in patent claims being narrowed, invalidated or held unenforceable, which could limit our ability to stop others from using or commercializing similar or identical technology and products or limit the duration of the patent protection of our technology and product candidates. Such proceedings also may result in substantial cost and require significant time from our scientists and management, even if the eventual outcome is favorable to us. Moreover, there could be public announcements of the results of hearings, motions or other developments related to any of the foregoing proceedings. If securities analysts or investors perceive those results to be negative, it could cause the price of shares of our Common Stock to decline. Any of the foregoing could harm our business, financial condition, results of operations and prospects. Moreover, some of our owned or in-licensed patents and patent applications may in the future be co-owned with third parties. If we are unable to obtain an exclusive license to any such co-owners' interest in such patents or patent applications, such co-owners may be able to license their rights to other third parties, including our competitors, who could market competing products and technology. In addition, we may need the cooperation of any such co-owners in order to enforce such patents against third parties, and such cooperation may not be provided to us. Our reliance on third parties requires us to share our trade secrets, which increases the possibility that a competitor will discover them or that our trade secrets will be misappropriated or disclosed. Because we rely on third parties to discover, develop and manufacture our product candidates, we must, at times, share certain of our trade secrets with them. We seek to protect our proprietary technology in part by entering into agreements containing confidentiality provisions, including if applicable, confidentiality agreements, material transfer agreements, collaborative research agreements, consulting agreements or other similar agreements with our collaborators, advisors, employees and consultants prior to beginning research or disclosing proprietary information. These agreements typically limit the rights of the third parties to use or disclose our confidential information, such as trade secrets. Despite these agreements with third parties, sharing trade secrets and other confidential information increases the risk that such trade secrets become known by our competitors, are inadvertently incorporated into the technology of others or are disclosed or used in violation of these agreements. Given that our proprietary position is based, in part, on our know-how and trade secrets, a competitor's discovery of our trade secrets or other unauthorized use or disclosure could impair our competitive position and may harm our business. In addition, these agreements typically restrict the ability of our advisors, employees, third-party contractors and consultants to publish data potentially relating to our trade secrets, although our agreements may contain certain limited publication rights. Despite our efforts to protect our trade secrets, our competitors may discover our trade secrets, either through breach of our agreements with third parties, independent development or publication of information by any of our third-party collaborators. A competitor's discovery of our trade secrets could impair our competitive position and have an adverse impact on our business. We may become involved in lawsuits to protect or enforce our patents or other intellectual property, which could be expensive, time-consuming and unsuccessful, and issued patents covering our technology and product candidates could be found invalid or unenforceable if challenged. Competitors and other third parties may infringe, misappropriate or otherwise violate our issued patents or other intellectual property or the patents or other intellectual property of our licensors. In addition, our patents or the patents of our licensors may become involved in inventorship or priority disputes. Our pending patent applications cannot be enforced against third parties practicing the technology claimed in such applications unless and until a patent issues from such applications. To counter infringement or other unauthorized use, we may be required to file infringement claims, which can be expensive and time-consuming. Any claims we assert against perceived infringers could provoke these parties to assert counterclaims against us alleging that we infringe their patents or that our patents are invalid and/or unenforceable. In a patent infringement proceeding, a court may decide that a patent of ours is invalid and/or unenforceable, in whole or in part, construe the patent's claims narrowly or refuse to stop the other party from using the technology at issue on the grounds that our patents do not cover the technology. An adverse result in any litigation proceeding could put one or more of our owned or licensed patents at risk of being invalidated and/or held unenforceable, interpreted narrowly or interpreted in a manner that would not prevent competitors from entering the market. Further, we may find it impractical or undesirable to enforce our intellectual property against some third parties. In patent litigation in the U.S., defendant counterclaims alleging invalidity and/or unenforceability are commonplace. Grounds for a validity challenge could be an alleged failure to meet any of several statutory requirements, including, e.g., lack of novelty, obviousness, non-enablement or insufficient written description. Grounds for an unenforceability assertion could be an allegation that someone connected with prosecution of the patent withheld relevant information from the USPTO or made a misleading statement during prosecution. Third parties may also raise similar claims before the USPTO or an equivalent foreign body, even outside the context of litigation. Potential proceedings include re-examination, post-grant review, inter partes review, interference proceedings, derivation proceedings and equivalent proceedings in foreign jurisdictions (e.g., opposition proceedings). Such proceedings could result in the revocation of, cancellation of, or amendment to our patents in such a way that they no longer cover our technology or any product candidates that we may develop. The outcome following legal assertions of invalidity and unenforceability is unpredictable. With respect to the validity question, for example, we cannot be certain that there is no invalidating prior art of which we and the patent examiner were unaware during prosecution. If a defendant were to prevail on a legal assertion of invalidity and/or unenforceability, we would lose at least part, and perhaps all, of the patent protection on the applicable product candidates or technology covered by the patent rendered invalid and/or unenforceable. Such a loss of patent protection could materially harm our business, financial condition, results of operations and prospects. Interference or derivation proceedings provoked by third parties or brought by us or declared by the USPTO may be necessary to determine the ownership or priority of inventions with respect to our patents or patent applications. An unfavorable outcome could require us to cease using the related technology or to attempt to license rights to it from the prevailing party. Such licenses may not be available on commercially reasonable terms, or at all, or may be non-exclusive. If we are unable to obtain and maintain such licenses, we may need to cease the development, manufacture and commercialization of one or more of the product candidates we may develop. In addition, if we or our licensors are unsuccessful in any inventorship disputes to which we or they are subject, we may lose valuable intellectual property rights, such as exclusive ownership of, or the exclusive right to use, our owned or in-licensed patents. The loss of exclusivity or the narrowing of scope of our owned and/or licensed patents could limit our ability to stop others from using or commercializing similar or identical technology and products. Any of the foregoing could result in a material adverse effect on our business, financial condition, results of operations or prospects. Even if we are successful in any of the foregoing disputes, it could result in substantial costs and be a distraction to management and other employees. Furthermore, because of the substantial amount of discovery required in connection with intellectual property litigation, there is a risk that some of our confidential information could be compromised by disclosure during this type of litigation or proceeding. Such litigation or proceedings could substantially increase our operating losses and reduce the resources available for development activities or any future sales, marketing or distribution activities. We may not have sufficient financial or other resources to conduct such litigation or proceedings adequately. Most of our competitors are larger than we are and have substantially greater resources. They are, therefore, more likely to be able to sustain the costs of complex patent litigation or proceedings than we can because of their greater financial resources and more mature and developed intellectual property portfolios. Accordingly, despite our efforts, we may not be able to prevent third parties from infringing, misappropriating or otherwise violating our intellectual property. Even if 55 Å resolved in our favor, litigation or other legal proceedings relating to intellectual property claims could result in substantial costs and diversion of management resources, which could harm our business. In addition, the uncertainties associated with litigation could compromise our ability to raise the funds necessary to continue our clinical trials, continue our internal research programs or in-license needed technology or other product candidates. There could also be public announcements of the results of the hearing, motions or other interim proceedings or developments. If securities analysts or investors perceive those results to be negative, it could cause the price of shares of our Common Stock to decline. Any of the foregoing events could harm our business, financial condition, results of operations and prospects. We may not be able to protect our intellectual property rights throughout the world. Filing, prosecuting, maintaining, defending and enforcing patents and other intellectual property rights on our product candidates in all countries throughout the world would be prohibitively expensive, and our intellectual property rights in some countries outside the U.S. can be less extensive than those in the U.S. As such, we may choose not to seek to protect our intellectual property in certain jurisdictions, which could leave us without recourse to prevent competitive products from being manufactured or commercialized in such jurisdictions. In addition, the laws of some foreign countries do not protect intellectual property rights to the same extent as federal and state laws in the U.S. Consequently, we may not be able to prevent third parties from practicing our inventions in all countries or from selling or importing products made using our inventions in all jurisdictions. Competitors may use our technologies in jurisdictions where we have not obtained patent protection or other intellectual property rights to develop their own products and may export otherwise infringing, misappropriating or violating products to territories where we have patent or other intellectual property protection, but enforcement rights are not as strong as those in the U.S. These products may compete with our product candidates, and our patents or other intellectual property rights may not be effective or sufficient to prevent them from competing. Many companies have encountered significant problems in protecting and defending intellectual property rights in foreign jurisdictions. The legal systems of some countries do not favor the enforcement of patents and other intellectual property rights, which could make it difficult for us to stop the infringement, misappropriation or other violation of our intellectual property rights generally. Proceedings to enforce our intellectual property rights in foreign jurisdictions could result in substantial costs and divert our efforts and attention from other aspects of our business, could put our patents at risk of being invalidated or interpreted narrowly and our patent applications at risk of not issuing and could provoke third parties to assert claims against us. We may not prevail in any lawsuits that we initiate, and the damages or other remedies awarded, if any, may not be commercially meaningful. Many countries, including European Union countries, India, Japan and China, have compulsory licensing laws under which a patent owner may be compelled under specified circumstances to grant licenses to third parties. In addition, many countries limit the enforceability of patents against government agencies or government contractors. In those countries, we may have limited remedies if patents are infringed or if we are compelled to grant a license to a third party, which could materially diminish the value of those patents. This could limit our potential revenue opportunities. Accordingly, our efforts to enforce our intellectual property rights around the world may be inadequate to obtain a significant commercial advantage from the intellectual property that we develop or license, which could adversely affect our business, financial condition, results of operations and prospects. We may not identify relevant third-party patents or pending patent applications or may incorrectly interpret the relevance, scope or expiration of a third-party patent which might adversely affect our ability to develop and market our product candidates. We are developing certain product candidates in highly competitive areas and cannot guarantee that any patent searches or analyses that we may conduct, including the identification of relevant patents or pending patent applications, the scope of patent claims or the expiration of relevant patents, are complete or thorough, nor can we be certain that we have identified each and every third-party patent and pending patent application in the U.S. and abroad that is or may be relevant to or necessary for the commercialization of our product candidates in any jurisdiction. For example, U.S. patent applications filed before November 29, 2000 and certain U.S. patent applications filed after that date that will not be filed outside the U.S. remain confidential until patents issue. Patent applications in the U.S. and elsewhere are published approximately 18 months after the earliest filing for which priority is claimed, with such earliest filing date being commonly referred to as the priority date. Therefore, patents or pending patent applications covering our product candidates could have been or may be filed in the future by third parties without our knowledge. Additionally, patents and pending patent applications that have been published can, subject to certain limitations, be later amended in a manner that could cover our product candidates or the manufacturing or use of our product candidates. The scope of a patent claim is determined by an interpretation of the law, the written disclosure in a patent and the patent's prosecution history. Our interpretation of the relevance or the scope of a patent or a pending patent application may be incorrect, which may negatively impact our ability to market our product candidates. We may incorrectly determine that our product candidates are not covered by a third-party patent or pending patent application or may incorrectly predict whether a third party's pending

application will issue with claims of relevant scope. Our determination of the expiration date of any patent in the U.S. or abroad that we consider relevant may be incorrect, which may negatively impact our ability to develop and market our product candidates. Our failure to identify and correctly interpret relevant patents or pending patent applications may negatively impact our ability to develop and market our product candidates. A 56 Å If we fail to identify or correctly interpret relevant patents or pending patent applications or if we are unable to obtain licenses to relevant patents or pending patent applications, we may be subject to infringement claims. We cannot guarantee that we will be able to successfully settle or otherwise resolve such infringement claims. If we fail in any such dispute, in addition to being forced to pay damages, potentially including in the form of future royalties, which may be significant, we may be temporarily or permanently prohibited from commercializing any of our product candidates that are held to be infringing. We might, if possible, also be forced to redesign product candidates so that we no longer infringe the third-party intellectual property rights. Any of these events, even if we were ultimately to prevail, could require us to divert substantial financial and management resources that we would otherwise be able to devote to our business and could adversely affect our business, financial condition, results of operations and prospects. If we are unable to obtain licenses from third parties on commercially reasonable terms or fail to comply with our obligations under such agreements, our business could be harmed. It may be necessary for us to use the patented or other proprietary technology of third parties to commercialize our products, in which case we would be required to obtain a license or ownership from these third parties. The licensing or acquisition of third-party intellectual property rights is a competitive area, and more established companies may pursue strategies to license or acquire third-party intellectual property rights that we may consider attractive or necessary. These established companies may have a competitive advantage over us due to their size, capital resources or greater clinical development and commercialization capabilities. In addition, companies that perceive us to be a competitor may be unwilling to assign or license rights to us. If we are unable to license or acquire such intellectual property or technology, or if we are forced to in-license such intellectual property or technology on unfavorable terms, our business could be materially harmed. If we are unable to obtain a necessary license, we may be unable to develop or commercialize the affected product candidates, or the cost of development, manufacture or commercialization may be materially increased, which could materially harm our business, and the third parties owning such intellectual property rights could seek either an injunction prohibiting our sales, or, with respect to our sales, an obligation on our part to pay royalties and/or other forms of compensation. Even if we are able to obtain a license, it may be non-exclusive, thereby giving our competitors access to the same technologies licensed to us. If we fail to comply with our obligations under any future license agreements, such counterparties may have the right to terminate these agreements, in which event we might not be able to develop, manufacture or commercialize, or may be forced to cease developing, manufacturing or marketing, any product that is covered by these agreements or may face other penalties under such agreements. Such an occurrence could materially adversely affect the value of the product candidate being developed under any such agreement. Termination of these agreements or reduction or elimination of our rights under these agreements may result in our having to negotiate new or reinstated agreements with less favorable terms, cause us to lose our rights under these agreements, including our rights to important intellectual property or technology, or impede, delay or prohibit the further development or commercialization of one or more product candidates that rely on such agreements. If we were to lose our rights to licensed intellectual property, we may not be able to continue developing or commercializing our product candidates, if approved. If we breach any of the agreements under which we license the use, development and commercialization rights to our product candidates or technology from third parties or, in certain cases, we fail to meet certain development deadlines, we could lose license rights that are important to our business. Patent terms may be inadequate to protect our competitive position on our product candidates for an adequate amount of time. Patents have a limited lifespan. In the U.S., if all maintenance fees are timely paid, the natural expiration of a patent is generally 20 years from its earliest U.S. non-provisional filing date. Various extensions may be available, and a given patent may be subject to other term adjustments, but the life of a patent, and the protection it affords, is limited. Even if patents covering our product candidates are obtained, once the patent life has expired for a product candidate, we may be open to competition from competitive products, including generic medications. Given the amount of time required for the development, testing and regulatory review of new product candidates, patents protecting such product candidates might expire before or shortly after such product candidates are commercialized. As a result, our owned and licensed patent portfolio may not provide us with sufficient rights to exclude others from commercializing product candidates similar or identical to ours. Depending upon the timing, duration and conditions of any FDA marketing approval of our product candidates, one or more of our U.S. patents may be eligible for limited patent term extension under the Drug Price Competition and Patent Term Restoration Act of 1984, referred to as the Hatch-Waxman Amendments, and one or more of our foreign patents may be eligible for patent term extension under similar legislation, for example, in the European Union. In the U.S., the Hatch-Waxman Amendments permit a patent term extension of up to five years for a patent covering an approved product as compensation for effective patent term lost during product development and the FDA regulatory review process provided other requirements are met. However, there are no assurances that the FDA, USPTO or any comparable foreign regulatory authority or national patent office will grant such extensions, in whole or in part and the length of any available extension may vary based on a number of factors. For example, we may not receive an extension if we fail to exercise due diligence during the testing phase or regulatory review process, fail to apply within applicable deadlines, fail to apply prior to expiration of relevant patents or otherwise fail to satisfy applicable requirements. Moreover, the length of the extension could be less than we request. Only one patent per approved product can be extended, the extension cannot extend the total patent term beyond 14 years from approval, and only those claims covering the approved drug, a method for using it or a method for manufacturing it may be extended. If we are unable to obtain patent term extension or the term of any such extension is less than we request, the period during Å 57 Å which we can enforce our patent rights for the applicable product candidate will be shortened, and our competitors may obtain approval to market competing products sooner. As a result, our revenue from applicable products could be reduced. Further, if this occurs, our competitors may take advantage of our investment in development and trials by referencing our clinical and preclinical data and launch their product earlier than might otherwise be the case, and our competitive position, business, financial condition, results of operations and prospects could be adversely affected. Changes in patent law could diminish the value of patents in general, thereby impairing our ability to protect our product candidates. Obtaining and enforcing patents in the pharmaceutical industry is inherently uncertain, due in part to ongoing changes in the patent laws. Depending on decisions by Congress, the federal courts, and the USPTO and equivalent institutions in other jurisdictions, the laws and regulations governing patents, and interpretation thereof, could change in unpredictable ways that could weaken our and our licensors' or collaborators' ability to obtain new patents or to enforce existing or future patents. For example, in recent years the U.S. Supreme Court has ruled on several patent cases that have been interpreted to have either narrowed the scope of patent protection or weakened the rights of patent owners in certain situations. Therefore, there is increased uncertainty with regard to our and our licensors' or collaborators' ability to obtain patents in the future, as well as uncertainty with respect to the value of patents once obtained. Recent patent reform legislation could increase the uncertainties and costs surrounding the prosecution of our and our licensors' or collaborators' patent applications and the enforcement or defense of our or our licensors' or collaborators' issued patents. Assuming that other requirements for patentability are met, prior to March 16, 2013, in the U.S., the first to invent the claimed invention was entitled to the patent, while outside the U.S., the first to file a patent application was entitled to the patent. On March 16, 2013, under the Leahy-Smith America Invents Act enacted in September 2011 (the "Leahy-Smith Act"), the U.S. transitioned to a first inventor to file system in which, assuming that other requirements for patentability are met, the first inventor to file a patent application will be entitled to the patent on an invention regardless of whether a third party was the first to invent the claimed invention. The Leahy-Smith Act also includes a number of significant changes that affect the way patent applications are prosecuted and may also affect patent litigation. These include allowing third-party submission of prior art to the USPTO during patent prosecution and additional procedures to attack the validity of a patent by USPTO-administered post-grant proceedings, including post-grant review, inter partes review and derivation proceedings. The USPTO recently developed new regulations and procedures to govern administration of the Leahy-Smith Act, and many of the substantive changes to patent law associated with the Leahy-Smith Act, particularly the first inventor-to-file provisions. Accordingly, it is not clear what, if any, impact the Leahy-Smith Act will have on the operation of our business. However, the Leahy-Smith Act and its implementation could increase the uncertainties and costs surrounding the prosecution of our or our licensors' patent applications and the enforcement or defense of our or our licensors' issued patents, all of which could harm our business, financial condition, results of operations and prospects. Obtaining and maintaining patent protection depends on compliance with various procedural, document submission, fee payment and other requirements imposed by governmental patent agencies, and our patent protection could be reduced or eliminated if we fail to comply with these requirements. Periodic maintenance fees, renewal fees, annuity fees and various other government fees on any issued patents and certain pending patent applications are required to be paid to the USPTO or foreign patent agencies in several stages over the lifetime of a patent. In certain circumstances, we may rely on our licensors to pay these fees. The USPTO and various foreign patent agencies also require compliance with a number of procedural, documentary, fee payment and other similar requirements during the patent application and prosecution process. Noncompliance events that could result in abandonment or lapse of a patent or patent application include failure to respond to official communications within prescribed time limits, non-payment of fees and failure to properly legalize and submit formal documents. While an inadvertent lapse can in many cases be cured by payment of a late fee or by other means in accordance with the applicable rules, there are situations in which non-compliance can result in irrevocable abandonment or lapse of the patent or patent application, resulting in partial or complete loss of patent rights in the relevant jurisdiction. If we or our licensors or collaborators fail to maintain the patents and patent applications covering our product candidates, our competitors might be able to enter the market with similar or identical products or technology, which would harm our business, financial condition, results of operations and prospects. Third parties may initiate legal proceedings alleging that we are infringing, misappropriating or otherwise violating their intellectual property rights, the outcome of which would be uncertain and could negatively impact the success of our business. Our commercial success depends upon our ability and the ability of our collaborators to develop, manufacture, market and sell our product candidates and use our proprietary technologies without infringing, misappropriating or otherwise violating the intellectual property and other proprietary rights of third parties. There is considerable intellectual property litigation in the biotechnology and pharmaceutical industries. We may become party to, or be threatened with, future adversarial proceedings or litigation regarding intellectual property rights with respect to our product candidates and technology, including re-examination, interference, post-grant review, inter partes review or derivation proceedings, or other similar proceedings, before the USPTO, a federal court or an equivalent foreign body. Numerous U.S. and foreign issued patents and pending patent applications owned by third parties exist in the fields in which we are developing our product candidates. In the event that any of these patents were asserted against us, we believe that we would have defenses against any such action, including that such patents are invalid and/or unenforceable, that our product candidates Å 58 Å do not infringe such patents, or that we would be able to replace such technology with alternative, non-infringing technology. However, if any such patents were to be asserted against us and our defenses to such assertion were unsuccessful and such alternative technology was not available or technologically or commercially practical, unless we obtain a license to such patents, we could be liable for damages, which could be significant and include treble damages and attorneys' fees if we are found to willfully infringe such patents, and we could be precluded from commercializing any product candidates that were ultimately held to infringe such patents. Any potential future legal proceedings relating to these patents could cause us to incur significant expenses, and could distract our technical and management personnel from their normal responsibilities. If we are unsuccessful in our challenges to these patents and become subject to litigation or are unable to obtain a license on commercially reasonable terms with respect to these patents, it could harm our business, financial condition, results of operations and prospects. Third parties may assert infringement claims against us based on existing patents or patents that may be granted in the future, regardless of their merit. Even if we believe third-party intellectual property claims are without merit, there is no assurance that a court would find in our favor on questions of infringement, validity, enforceability or priority. A court of competent jurisdiction could hold that third-party patents asserted against us are valid, enforceable and infringed, which could materially and adversely affect our ability to commercialize any product candidates we may develop and any other product candidates or technologies covered by the asserted third-party patents. In order to successfully challenge the validity of any such U.S. patent in federal court, we would need to overcome a presumption of validity. As this burden is a high one requiring us to present clear and convincing evidence as to the invalidity of any such U.S. patent claim, there is no assurance that a court of competent jurisdiction would invalidate the claims of any such U.S. patent. If we are found to infringe, misappropriate or otherwise violate a third party's intellectual property rights, and we are unsuccessful in demonstrating that such rights are invalid or unenforceable, we could be required to obtain a license from such a third party in order to continue developing and marketing our products and technology. However, we may not be able to obtain any required license on commercially reasonable terms or at all. Even if we were able to obtain a license, it could be non-exclusive, thereby giving our competitors access to the same technologies licensed to us, and it could require us to make substantial licensing and royalty payments. We could be forced, including by court order, to cease commercializing the infringing technology or product candidates. A finding of infringement could prevent us from commercializing our product candidates or force us to cease some of our business operations. In the event of a successful claim of infringement against us, we may have to pay substantial damages, including treble damages and attorneys' fees for willful infringement, pay royalties and other fees, redesign our infringing drug or obtain one or more licenses from third parties, which may be impossible or require substantial time and monetary expenditure. Claims that we have misappropriated the confidential information or trade secrets of third parties could have a similar negative impact on our business. Any of the foregoing events would harm our business, financial condition, results of operations and prospects. We may be subject to claims by third parties asserting that we or our employees have infringed upon, misappropriated or otherwise violated their intellectual property rights, or claiming ownership of what we regard as our own intellectual property. Many of our employees were previously employed at other biotechnology or pharmaceutical companies. Although we try to ensure that our employees, consultants and advisors do not use the proprietary information or know-how of others in their work for us, we may be subject to claims that we or these individuals have used or disclosed intellectual property, including trade secrets or other proprietary information, of others, such as any such individual's former employer. Litigation may be necessary to defend against these claims. In addition, we or our licensors may be subject to claims that former employees, collaborators or other third parties have an interest in our owned or in-licensed patents or other intellectual property as an inventor or co-inventor. While it is our policy to require our employees and contractors who may be involved in the development of intellectual property to execute agreements assigning such intellectual property to us, we may be unsuccessful in executing such an agreement with each party who in fact conceives, develops or reduces to practice intellectual property that we regard as our own. Our and their assignment agreements may not be self-executing or may be breached, and we may be forced to bring claims against third parties, or defend claims they may bring against us, to determine the ownership of what we regard as our intellectual property. If we fail in prosecuting or defending any such claims, in addition to paying monetary damages, we may lose valuable intellectual property rights or personnel. Even if we are successful in litigating such claims, litigation could result in substantial costs, delay development of our product candidates and be a distraction to management. Any of the foregoing events would harm our business, financial condition, results of operations and prospects. Intellectual property litigation could cause us to spend substantial resources and distract our personnel from their normal responsibilities. Even if resolved in our favor, litigation or other legal proceedings relating to intellectual property claims may cause us to incur significant expenses and could distract our technical and management personnel from their normal responsibilities. In addition, there could be public announcements of the results of hearings, motions or other interim proceedings or developments and if securities analysts or investors perceive these results to be negative, it could have a substantial adverse effect on the price of our Common Stock. Such litigation or proceedings could substantially increase our operating losses and reduce the resources available for development activities Å 59 Å or any future sales, marketing or distribution activities. We may not have sufficient financial or other resources to conduct such litigation or proceedings adequately. Some of our competitors may be able to sustain the costs of such litigation or proceedings more effectively than we can because of their greater financial resources. Uncertainties resulting from the initiation and continuation of patent litigation or other proceedings could compromise our ability to compete in the marketplace, including compromising our ability to raise the funds necessary to continue our clinical trials, continue our research programs, license necessary technology from third parties or enter into development collaborations that would help us commercialize our product candidates, if approved. Any of the foregoing events would harm our business, financial condition, results of operations and prospects. If we are unable to protect the confidentiality of our trade secrets, our business and competitive position would be harmed. We rely on trade secrets and agreements containing confidentiality obligations to protect our unpatented know-how, technology and other proprietary information and to maintain our competitive position. With respect to our research and development programs, we consider trade secrets and know-how to be one of our important sources of intellectual property, including our extensive knowledge of certain drug delivery techniques and drug conjugation. Trade secrets and know-how can be difficult to protect. We seek to

protect these trade secrets and other proprietary technology, in part, by entering into non-disclosure and confidentiality agreements with parties who have access to them, such as our employees, collaborators, CROs, contract manufacturers, consultants, advisors and other third parties. We also enter into confidentiality and invention or patent assignment agreements with our employees and consultants. We cannot guarantee that we have entered into such agreements with each party that may have or has had access to our trade secrets or proprietary technology and processes. Despite these efforts, any of these parties may breach the agreements and disclose our proprietary information, including our trade secrets, and we may not be able to obtain adequate remedies for such breaches. Enforcing a claim that a party illegally disclosed or misappropriated a trade secret is difficult, expensive and time-consuming, and the outcome is unpredictable. In addition, some courts inside and outside the U.S. are less willing or unwilling to protect trade secrets. If any of our trade secrets were to be lawfully obtained or independently developed by a competitor or other third party, we would have no right to prevent them from using that technology or information to compete with us. If any of our trade secrets were to be disclosed to or independently developed by a competitor or other third party, our competitive position would be materially and adversely harmed. We may not be able to protect and enforce our trademarks and trade names, or build name recognition in our markets of interest thereby harming our competitive position. We intend to rely on both registered and common law rights for our trademarks. We have applied to register certain of our trademarks with the USPTO and trademark authorities in certain other countries and may in the future seek to register additional trademarks in the U.S. or other countries. Our current and future trademark applications may not mature to registration in a timely fashion or at all, and our registered trademarks may not be maintained or enforced. In the U.S. and some foreign jurisdictions, our ability to obtain and maintain trademark registrations and acquire enforceable trademark rights depends on making use of our marks in commerce, meaning we must make a certain amount of progress, depending on the jurisdiction, in our clinical studies or in the commercialization of our products. If we fail to satisfy these requirements or any other requirements of applicable regulatory authorities, we may not have enforceable trademark rights or registrations in such jurisdictions. We have yet to obtain trademark registrations for the NUATION or NUATION BIO trademarks in the U.S., and we have yet to apply to register any brand name for any product candidate in the U.S. or any other jurisdiction. In addition, the registered or unregistered trademarks or trade names that we own may be challenged, infringed, circumvented, declared generic, lapsed or determined to be infringing on or dilutive of other marks. We may be unable to develop any enforceable trademark rights in relevant countries, or to protect the rights that we do develop. We may be forced to stop using our trademarks or trade names, which we need for name recognition by potential partners and customers in our markets of interest, and spend time and money rebranding. In addition, third parties have filed, and may in the future file, for registration of trademarks similar or identical to our trademarks, thereby impeding our ability to build brand identity and possibly leading to market confusion. If they succeed in registering or developing common law rights in such trademarks, and if we are not successful in enforcing our rights, we may not be able to use these trademarks to develop brand recognition of our company, technologies, products or services. In addition, there could be potential trade name or trademark infringement litigation brought against us by owners of other trademarks that incorporate variations of our registered or unregistered trademarks or trade names. During the trademark registration process, we may receive office actions from the USPTO or from comparable agencies in foreign jurisdictions refusing to register our trademarks. Although we would be given an opportunity to respond to those refusals, we may be unable to overcome them. In addition, in the USPTO and in comparable agencies in many foreign jurisdictions, third parties are given an opportunity to oppose pending trademark applications and to seek the cancellation of registered trademarks. Opposition or cancellation proceedings may in the future be filed against our trademark applications or registrations, and our trademark applications or registrations may not survive such proceedings. In addition, third parties may file first for our trademarks or similar variations thereof in certain countries. If they succeed in registering such trademarks, and if we are not successful in challenging such third-party rights, we may not be able to use these trademarks to market our products in those countries. If we do not secure registrations for our trademarks, we may encounter more difficulty in enforcing them against third parties than we otherwise would. If we are unable to establish name recognition based on our trademarks and trade names, we may be unable to compete effectively, which could have an adverse effect on our business, financial condition, results of operations and prospects. Intellectual property rights do not necessarily address all potential threats. The degree of future protection afforded by our intellectual property rights is uncertain because intellectual property rights have limitations and may not adequately protect our business or permit us to maintain our competitive advantage. For example, others may be able to make products that are similar to any product candidates we may develop or utilize similar technology but that are not covered by the claims of the patents that we own or license now or in the future; we, or our current or future licensors, might not have been the first to make the inventions covered by the issued patents or pending patent applications that we own or license now or in the future; we, or our current or future licensors, might not have been the first to file patent applications covering certain of our or their inventions; others may independently develop similar or alternative technologies or duplicate any of our technologies without infringing our owned or licensed intellectual property rights; it is possible that our pending owned or licensed patent applications or those that we may own or license in the future will not lead to issued patents; issued patents that we hold rights to may be held invalid or unenforceable, including as a result of legal challenges by other persons; our competitors might conduct research and development activities in the U.S. under FDA-related safe harbor patent infringement exemptions and/or in countries where we do not have patent rights and then use the information learned from such activities to develop competitive products for sale in our major commercial markets; we may not develop additional proprietary technologies that are patentable; the patents or pending patent applications of others may harm our business; and we may choose not to file for patent protection in order to maintain certain trade secrets or know-how, and a third party may subsequently file for and obtain a patent covering such intellectual property. Should any of these events occur, they could harm our business, financial condition, results of operations and prospects. Risks Related to Our Business Operations, Employee Matters and Managing Growth AnHeart could be difficult to integrate, divert the attention of management, and disrupt our business, and the anticipated synergies and other benefits of the AnHeart Acquisition may not be realized in the amounts anticipated, or may not be realized within the anticipated timeframe, or at all. It is possible that there could be a loss of our existing or AnHeart's key employees and customers, disruption of either company's or both companies' ongoing businesses or unexpected issues, higher than expected costs and an overall post-completion process that takes longer than originally anticipated. Specifically, the following issues, among others, must be addressed in combining AnHeart's operations with ours in order to realize the anticipated benefits of the AnHeart Acquisition so the combined company performs as the parties hope: combining the companies' corporate functions; combining their business with our business in a manner that permits us to achieve the synergies anticipated to result from the Acquisition, the failure of which would result in the anticipated benefits of the Acquisition not being realized in the time frame currently anticipated or at all; maintaining existing and new agreements with customers, service providers, and vendors; determining whether and how to address possible differences in corporate cultures, management philosophies and strategies relating to channels, resellers, and partners; integrating the companies' administrative and information technology infrastructure; developing products and technology that allow value to be unlocked in the future; and evaluating and forecasting the financial impact of the Acquisition transaction, including accounting impacts. Failure to address any of the above-listed issues could have a material adverse effect on our business, results of operations and financial position. In addition, at times the attention of certain members of our management and resources may be focused on integration planning of the businesses of the two companies and diverted from day-to-day business operations, which may disrupt our ongoing business and the business of the combined company. The failure to meet the challenges involved in combining the two companies could, among other things, cause an interruption of, or a loss of momentum in, our activities and could adversely affect our results of operations. The overall combination of the two companies may also result in, among other things, material unanticipated problems, expenses, liabilities, competitive responses and loss of customer and other business relationships. The difficulties of combining our operations include, among others: diversion of management and employee attention to integration matters; difficulties in integrating operations and systems, including, but not limited to, communications systems, administrative and information technology infrastructure, financial reporting and internal control systems; challenges in conforming standards, controls, procedures and accounting and other policies, business cultures and compensation structures between the two companies; difficulties in integrating employees and teams of the respective businesses and attracting and retaining key personnel; challenges in retaining and obtaining customers, suppliers and other commercial relationships; difficulties in managing the expanded operations of a larger and more complex company; and potential unknown liabilities, adverse consequences and unforeseen increased expenses associated with the integration. Many of these factors are outside of our control and any of them could result in lower revenues, higher costs and diversion of management time and energy, which could materially impact our business, financial condition and results of operations. In addition, even if the operations of the companies are integrated successfully, the full benefits of the AnHeart Acquisition may not be realized, including, among others, the synergies, cost savings or sales or growth opportunities that are expected. These benefits may not be achieved within the anticipated time frame or at all. As a result, it cannot be assured that the integration will result in the realization of the full benefits expected from the AnHeart Acquisition within the anticipated time frames, or at all. Our business, operations and clinical development plans and timelines and supply chain could be adversely affected by the effects of health epidemics on the manufacturing, clinical trial and other business activities performed by us or by third parties with whom we conduct business, including our CMOs, CROs, shippers and others. Our business could be adversely affected by health epidemics wherever we have clinical trial sites or other business operations. In addition, health epidemics could cause significant disruption in the operations of CMOs, CROs and other third parties upon whom we rely. For example, the COVID-19 pandemic presented a substantial public health and economic challenge around the world and affected employees, patients, communities and business operations, as well as the U.S. economy and financial markets. Geographic regions imposed shelter-in-place orders, quarantines or similar orders or restrictions to control the spread of epidemic disease. These and similar, and perhaps more severe, disruptions in our operations could negatively impact our business, operating results and financial condition. We are dependent on a worldwide supply chain for products to be used in our clinical trials and, if approved by the regulatory authorities, for commercialization. Quarantines, shelter-in-place and similar government orders, or the expectation that such orders, shutdowns or other restrictions could occur may impact personnel at third-party manufacturing facilities in the U.S. and other countries, or the availability or cost of materials or supplies, which could disrupt our supply chain or our ability to enroll patients in or perform testing for our clinical trials. In addition, closures of transportation carriers and modal hubs could materially impact our clinical development and any future commercialization timelines. If our relationships with our suppliers or other vendors are terminated or scaled back as a result of health epidemics, we may not be able to enter into arrangements with alternative suppliers or vendors or do so on commercially reasonable terms or in a timely manner. Switching or adding additional suppliers or vendors involves substantial cost and requires management time and focus. In addition, there is a natural transition period when a new supplier or vendor commences work. As a result, delays generally occur, which could adversely impact our ability to meet our desired clinical development and any future commercialization timelines. Although we carefully manage our relationships with our suppliers and vendors, there can be no assurance that we will not encounter challenges or delays in the future or that these delays or challenges will not have an adverse impact on our business, financial condition and prospects. See Risk Factors "Risks Related to Our Dependence on Third Parties." A62 In addition, our clinical trials may be affected by health epidemics. In the future, clinical site initiation and patient enrollment may be delayed due to prioritization of hospital resources toward such health epidemics or concerns among patients about participating in clinical trials during a health epidemic and public health measures imposed by the respective national governments of countries in which the clinical sites are located. Some patients may have difficulty following certain aspects of clinical trial protocols if quarantines impede patient movement or interrupt healthcare services. Similarly, our inability to successfully recruit and retain patients and principal investigators and site staff who, as healthcare providers, may have heightened exposure to epidemic disease or experience additional restrictions by their institutions, city or state governments could adversely impact our clinical trial operations. Health epidemics may lead to disruption and volatility in the global capital markets, which increases the cost of, and adversely impacts access to, capital and increases economic uncertainty. Health epidemics may also result in volatile trading prices for the Common Stock of biopharmaceutical companies. To the extent health epidemics adversely affect our business, financial results and value of our Common Stock, it may also affect our ability to access capital, which could in the future negatively affect our liquidity. Our future success depends on our ability to retain Dr. Hung and our other key employees, consultants and advisors and to attract, retain and motivate qualified personnel. We are highly dependent on the management, research and development, clinical, financial and business development expertise of Dr. Hung and our executive officers, as well as the other members of our scientific and clinical teams. Although we have employment offer letters with each of our executive officers, each of them may terminate their employment with us at any time. We do not maintain a key person life insurance for any of our executives or employees. Recruiting and retaining qualified scientific and clinical personnel and, if we are successful in obtaining marketing approval for our product candidates, sales and marketing personnel, is critical to our success. The loss of the services of our executive officers or other key employees could impede the achievement of our research, development and commercialization objectives and seriously harm our ability to successfully implement our business strategy. In particular, in light of Dr. Hung's central role in the discovery of all of our current product candidates, our ongoing discovery activities and development programs, the recruitment of our other executives and key employees and all other aspects of our strategy and operations, we believe our loss of Dr. Hung's services for any reason would severely impair our business and prospects. Replacing executive officers and key employees may be difficult and may take an extended period of time because of the limited number of individuals in our industry with the breadth of skills and experience required to successfully develop, gain regulatory approval for and commercialize our product candidates. Competition to hire qualified personnel in our industry is intense, and we may be unable to hire, train, retain or motivate these key personnel on acceptable terms given the competition among numerous pharmaceutical and biotechnology companies for similar personnel. Furthermore, to the extent we hire personnel from competitors, we may be subject to allegations that they have been improperly solicited or that they have divulged proprietary or other confidential information, or that their former employers own their research output. We also experience competition for the hiring of scientific and clinical personnel from universities and research institutions. In addition, we rely on consultants and advisors, including scientific and clinical advisors, to assist us in formulating our research and development and commercialization strategy. Our consultants and advisors may be employed by employers other than us and may have commitments under consulting or advisory contracts with other entities that may limit their availability to us. If we are unable to continue to attract and retain high quality personnel, our ability to pursue our growth strategy will be limited, and could harm our business, financial condition, results of operations and prospects. We expect to expand our development, regulatory, sales, marketing and distribution capabilities, and as a result, we may encounter difficulties in managing our growth, which could disrupt our operations. As of June 30, 2024, we had 167 employees. As our preclinical and clinical development progresses, we expect to experience growth in the number of our employees and the scope of our operations, particularly in the areas of research, clinical operations, regulatory affairs, general and administrative and, if any of our product candidates receives marketing approval, sales, marketing and distribution. To manage our anticipated future growth, we must continue to implement and improve our managerial, operational and financial systems, expand our facilities and continue to recruit and train additional qualified personnel. Due to our limited financial resources and the limited experience of our management team in managing a company with such anticipated growth, we may not be able to effectively manage the expansion of our operations or recruit and train additional qualified personnel. The expansion of our operations may lead to significant costs and may divert our management and business development resources. Any inability to manage growth could delay the execution of our business plans or disrupt our operations. Our employees, independent contractors, consultants, commercial collaborators, principal investigators, CROs and vendors may engage in misconduct or other improper activities, including non-compliance with regulatory standards and requirements. We are exposed to the risk that our employees, independent contractors, consultants, commercial collaborators, principal investigators, CROs and vendors may engage in fraudulent conduct or other illegal activity. Misconduct by these parties could include a63 intentional, reckless or negligent conduct or unauthorized activities that violates (1) the laws, regulations and guidance of the FDA, the European Commission, the EMA, the MHRA, NMPA and other similar regulatory authorities, including those laws requiring the reporting of true, complete and accurate information to such authorities, (2) manufacturing standards, (3) federal and state data privacy, security, fraud and abuse and other healthcare laws and regulations in the U.S. and abroad and (4) laws that require the true, complete and accurate reporting of financial information or data. In particular, sales, marketing and business arrangements in the healthcare industry are subject to extensive laws and regulations intended to prevent fraud, misconduct, kickbacks, self-dealing and other abusive practices. These laws and regulations may restrict or prohibit a wide range of pricing, discounting, marketing and promotion, sales commission, customer incentive programs and other business arrangements. Misconduct by these parties could also involve the improper use of individually identifiable information, including information obtained in the course of clinical trials, creating fraudulent data in our preclinical studies or clinical trials or illegal misappropriation of product candidates, which could result in regulatory sanctions and serious harm to our reputation. We

have adopted a code of business conduct and ethics, but it is not always possible to identify and deter misconduct by employees and other third parties, and the precautions we take to detect and prevent this activity may not be effective in controlling unknown or unmanaged risks or losses or in protecting us from governmental investigations or other actions or lawsuits stemming from a failure to be in compliance with such laws or regulations. Additionally, we are subject to the risk that a person or government could allege such fraud or other misconduct, even if none occurred. If any such actions are instituted against us, and we are not successful in defending ourselves or asserting our rights, those actions could have a significant impact on our business, including the imposition of significant civil, criminal and administrative penalties, including damages, fines, disgorgement, imprisonment, exclusion from participation in government healthcare programs, such as Medicare and Medicaid, contractual damages, reputational harm and the delay, reduction, termination or restructuring of our operations. If our information technology systems or those of third parties upon which we rely, or our data are or were compromised, we could experience adverse consequences resulting from such compromise, including but not limited to regulatory investigations or actions; litigation; fines and penalties; disruptions of our business operations; reputational harm; loss of revenue or profits; and other adverse consequences. In the ordinary course of business, we and the third parties with whom we work, process sensitive data, and, as a result, we and the third parties with whom we work face a variety of evolving threats that could cause security incidents. Cyber-attacks, malicious internet-based activity, online and offline fraud, and other similar activities threaten the confidentiality, integrity, and availability of our sensitive data and information technology systems, and those of the third parties upon with whom we work. Such threats are prevalent and continue to rise, are increasingly difficult to detect, and come from a variety of sources, including traditional computer hackers, threat actors, hacktivists, organized criminal threat actors, personnel (such as through theft or misuse), sophisticated nation states, and nation-state-supported actors. Some actors now engage and are expected to continue to engage in cyber-attacks, including without limitation nation-state actors for geopolitical reasons and in conjunction with military conflicts and defense activities. During times of war and other major conflicts, we and the third parties with whom we work may be vulnerable to a heightened risk of these attacks, including retaliatory cyber-attacks, that could materially disrupt our systems and operations, supply chain, and ability to produce, sell and distribute our services. We and the third parties upon which we rely are subject to a variety of evolving threats, including but not limited to social-engineering attacks (including through deep fakes, which may be increasingly more difficult to identify as fake, and phishing attacks), malicious code (such as viruses and worms), malware (including as a result of advanced persistent threat intrusions), denial-of-service attacks, credential stuffing, credential harvesting, personnel misconduct or error, ransomware attacks, supply-chain attacks, software bugs, server malfunctions, software or hardware failures, loss of data or other information technology assets, adware, attacks enhanced or facilitated by artificial intelligence (AI), telecommunications failures, earthquakes, fires, floods, and other similar threats. In particular, severe ransomware attacks are becoming increasingly prevalent and can lead to significant interruptions in our operations, ability to provide our products or services, loss of sensitive data and income, reputational harm, and diversion of funds. Extortion payments may alleviate the negative impact of a ransomware attack, but we may be unwilling or unable to make such payments due to, for example, applicable laws or regulations prohibiting such payments. Remote work has become more common and has increased risks to our information technology systems and data, as more of our personnel utilize network connections, computers, and devices outside our premises or network, including working at home, while in transit or in public locations. Additionally, future or past business transactions (such as acquisitions or integrations) could expose us to additional cybersecurity risks and vulnerabilities, as our systems could be negatively affected by vulnerabilities present in acquired or integrated entities' systems and technologies. Furthermore, we may discover security issues that we did not find during due diligence of such acquired or integrated entities, and it may be difficult to integrate companies into our information technology environment and security program. A 64 A In addition, our reliance on third parties could introduce new cybersecurity risks and vulnerabilities, including supply-chain attacks, and other threats to our business operations. We rely on various third parties and technologies to operate critical business systems to process sensitive data in a variety of contexts, including, without limitation, cloud-based infrastructure, data center facilities, encryption and authentication technology, personnel email, and other functions. We also rely on third parties to provide other products, services, parts, or otherwise to operate our business. Our ability to monitor these third parties' information security practices is limited, and these third parties may not have adequate information security measures in place. If the third parties with whom we work experience a security incident or other interruption, we could experience adverse consequences. While we may be entitled to damages if such third parties fail to satisfy their privacy or security-related obligations to us, any award may be insufficient to cover our damages, or we may be unable to recover such award. In addition, supply-chain attacks have increased in frequency and severity, and we cannot guarantee that third parties' infrastructure in our supply chain or our third-party partners' supply chains have not been compromised. While we have implemented security measures designed to protect against security incidents, there can be no assurance that these measures will be effective. We take steps designed to detect, mitigate, and remediate vulnerabilities in our information systems (such as our hardware and/or software, including that of third parties upon which we rely). We may not, however, detect and remediate all such vulnerabilities including on a timely basis. Further, we may experience delays in developing and deploying remedial measures and patches designed to address identified vulnerabilities. Vulnerabilities could be exploited and result in a security incident. Any of the previously identified or similar threats could cause a security incident or other interruption that could result in unauthorized, unlawful, or accidental acquisition, modification, destruction, loss, alteration, encryption, disclosure of, or access to our sensitive data or our information technology systems, or those of the third parties with whom we work. A security incident or other interruption could disrupt our ability (and that of third parties upon whom we rely) to provide our services. We may expend significant resources or modify our business activities (including our clinical trial activities) to try to protect against security incidents. Additionally, certain data privacy and security obligations may require us to implement and maintain specific security measures or industry-standard or reasonable security measures to protect our information technology systems and sensitive data. Applicable data privacy and security obligations may require us to notify relevant stakeholders, including affected individuals, regulators, and investors, of security incidents, or to implement other requirements, such as providing credit monitoring. Such disclosures and compliance with such requirements are costly, and the disclosure or the failure to comply with such requirements could lead to adverse consequences. If we (or a third party with whom we work) experience a security incident or are perceived to have experienced a security incident, we may experience adverse consequences, such as government enforcement actions (for example, investigations, fines, penalties, audits, and inspections); additional reporting requirements and/or oversight; restrictions on processing sensitive data (including personal data); litigation (including class claims); indemnification obligations; negative publicity; reputational harm; monetary fund diversions; diversion of management attention; interruptions in our operations (including availability of data); financial loss; and other similar harms. Security incidents and attendant consequences may negatively impact our ability to grow and operate our business. Our contracts may not contain limitations of liability, and even where they do, there can be no assurance that limitations of liability in our contracts are sufficient to protect us from liabilities, damages, or claims related to our data privacy and security obligations. We cannot be sure that our insurance coverage, if any, will be adequate or sufficient to protect us from or to mitigate liabilities arising out of our privacy and security practices, that such coverage will continue to be available on commercially reasonable terms or at all, or that such coverage will pay future claims. In addition to experiencing a security incident, third parties may gather, collect, or infer sensitive data about us from public sources, data brokers, or other means that reveals competitively sensitive details about our organization and could be used to undermine our competitive advantage or market position. Additionally, sensitive data of the Company could be leaked, disclosed, or revealed as a result of or in connection with our personnel's or vendors' use of generative AI technologies. Risks Related to Doing Business in China and Our International Operations Changes in the political and economic policies or in relations between China and the United States may affect our business, financial condition, and results of operations. Due to our operations in China, our business, results of operations, financial condition and prospects may be influenced to a certain degree by economic, political, legal and social conditions in China or changes in government relations between China and the United States or other governments. The Chinese government may intervene in or influence our operations, which could result in a change in our operations. Any economic downturn, whether actual or perceived, further decrease in economic growth rates or an otherwise uncertain economic outlook could affect our business, financial condition and results of operations. In addition, the global macroeconomic environment is facing challenges. It is unclear whether these challenges and uncertainties will be contained or resolved, and what effects they may have on the global political and economic conditions, and our business operations in the long term. There is a 65 A significant uncertainty about the future relationship between the United States and China with respect to trade policies, treaties, government regulations and tariffs. The Chinese government has implemented various measures to encourage economic development and guide the allocation of resources. Some of these measures may benefit the overall Chinese economy, but may have a negative effect on us. Due to our operations in China, any future Chinese, U.S. or other rules and regulations that place restrictions on capital raising or other activities by companies with operations in China could affect our business and results of operations. If the business environment in China deteriorates from the perspective of domestic or international investment, or if relations between China and the United States or other governments deteriorate and geopolitical tensions between China and the United States increase, our business in China and United States may also be affected. Changes in U.S. and Chinese regulations may impact our business, our operating results, and our ability to raise capital. The U.S. government, including the SEC, has made statements and taken certain actions that led to changes to United States and international relations, and will impact companies with connections to the United States or China, including imposing several rounds of tariffs affecting certain products manufactured in China, imposing certain sanctions and restrictions in relation to China and issuing statements indicating enhanced review of companies with certain operations based in China. It is unknown whether and to what extent new legislation, executive orders, tariffs, laws or regulations will be adopted, or the effect that any such actions would have on companies with significant connections to the United States or to China, our industry or on us. We conduct research activities and have business operations both in the United States and China. Any unfavorable government policies on cross-border relations and/or international trade, including increased scrutiny on companies with certain operations based in China, capital controls or tariffs, may affect the competitive position of our drug products, the hiring of scientists and other research and development personnel, the demand for our drug products, the import or export of raw materials in relation to drug development, our ability to raise capital, or prevent us from selling our drug products in certain countries. Furthermore, the SEC has issued statements primarily focused on companies with certain operations based in China, such as us. For example, in 2021, the Chairman of the SEC, issued a Statement on Investor Protection Related to Recent Developments in China, pursuant to which he stated that he has asked the SEC staff to engage in targeted additional reviews of filings for companies with certain operations based in China. The statement also addressed risks inherent in companies with variable interest entity (VIE) structures. We do not have a VIE structure and are not in an industry that is subject to foreign ownership limitations by China. However, it is possible that our periodic reports and other filings with the SEC may be subject to enhanced review by the SEC and this additional scrutiny could affect our ability to effectively raise capital in the United States. In response to the SEC's 2021 statement, the China Securities Regulatory Commission (CSRC) announced in August 2021, that it is our belief that Chinese and U.S. regulators shall continue to enhance communication with the principle of mutual respect and cooperation, and properly address the issues related to the supervision of China-based companies listed in the U.S. so as to form stable policy expectations and create benign rules framework for the market. While the CSRC will continue to collaborate closely with different stakeholders including investors, companies, and relevant authorities to further promote transparency and certainty of policies and implementing measures, it emphasized that it has always been open to companies' choices to list their securities on international or domestic markets in compliance with relevant laws and regulations. If any new legislation, executive orders, tariffs, laws and/or regulations are implemented, if existing trade agreements are renegotiated or if the U.S. or Chinese governments take retaliatory actions due to the recent U.S.-China tension, such changes could have an adverse effect on our business, financial condition and results of operations. Compliance with China's new Data Security Law, Cyber Security Law, Cybersecurity Review Measures, Personal Information Protection Law, regulations and guidelines relating to the multi-level protection scheme on cyber security and any other future laws and regulations may entail significant expenses and could affect our business. China has implemented or will implement rules and is considering a number of additional proposals relating to data protection. China's new Data Security Law took effect in September 2021. The Data Security Law provides that the data processing activities must be conducted based on data classification and hierarchical protection system for the purpose of data protection and prohibits entities in China from transferring data stored in China to foreign law enforcement agencies or judicial authorities without prior approval by the Chinese government. Additionally, China's Cyber Security Law, promulgated by the Standing Committee of the National People's Congress in 2016 and came into effect in 2017, and the Administrative Measures for the Hierarchical Protection of Information Security promulgated by the Ministry of Public Security, National Administration of State Secrets Protection, State Cryptography Administration and other government authority in 2007, requires companies to take certain organizational, technical and administrative measures and other necessary measures to ensure the security of their networks and data stored on their networks. Specifically, the Cyber Security Law provides that China adopt a multi-level protection scheme (MLPS), under which network operators are required to perform obligations of security protection to ensure that the network is free from interference, disruption or unauthorized access, and prevent network data from being disclosed, stolen or tampered. Under the MLPS, entities operating information systems must have a thorough assessment of the risks and the conditions of their information and network systems to determine the level of the entity's information and network systems. These levels range from the lowest Level 1 to the highest Level 5 pursuant to a series of national standards on the grading and implementation of the classified protection of cyber security. The grading result will determine the set of security protection obligations that entities must comply with. Entities classified as Level 2 or above should report the grade to the relevant government authority for examination and approval. In 2021, the Cyberspace Administration of China (CAC) published a draft revision to the existing Cybersecurity Review Measures for public comment (the Revised Draft CAC Measures). In 2022, together with 12 other Chinese regulatory authorities, the CAC released the final version of the Revised Draft CAC Measures (the Revised CAC Measures), which came into effect in 2022. Pursuant to the Revised CAC Measures, critical information infrastructure operators procuring network products and services, and online platform operators (as opposed to data processors in the Revised Draft CAC Measures) carrying out data processing activities which affect or may affect national security, shall conduct a cybersecurity review pursuant to the provisions therein. In addition, online platform operators possessing personal information of more than one million users seeking to be listed on foreign stock markets must apply for a cybersecurity review. In 2021, the CAC further published the Regulations on Network Data Security Management (Draft for Comment) (the Draft Management Regulations), under which data processors refer to individuals and organizations who determine the data processing activities in terms of the purpose and methods at their discretion. The Draft Management Regulations reiterate that data processors shall be subject to cybersecurity review if (i) they process personal information of more than one million persons and they are aiming to list on foreign stock markets, or (ii) their data processing activities affect or may affect Chinese national security. The Draft Management Regulations also request data processors seeking to list on foreign stock markets to annually assess their data security by themselves or through data security service organizations, and submit the assessment reports to relevant competent authorities. As the Draft Management Regulations are released only for public comment, the final version and the effective date thereof is subject to change. As of the date of this Report, we have not received any notice from any Chinese regulatory authority identifying us as a critical information infrastructure operator, an online platform operator or a data processor, or requiring us to go through the cybersecurity review procedures pursuant to the Revised CAC Measures and the Draft Management Regulations. Based on our understanding of the Revised CAC Measures, and the Draft Management Regulations if enacted as currently proposed, we do not expect to become subject to cybersecurity review by the CAC for issuing securities to foreign investors because: (i) the clinical and preclinical data we handle in our business operations, either by its nature or in scale, do not normally trigger significant concerns over PRC national security; and (ii) we have not processed, and do not anticipate to process in the foreseeable future, personal information for more than one million users or persons. However, there remains uncertainty as to how the Revised CAC Measures, and the Draft Management Regulations if enacted as currently proposed, will be interpreted or implemented; for example, neither the Revised CAC Measures nor the Draft Management Regulations provides further clarification or interpretation on the criteria for determining those activities that affect or may affect national security and relevant Chinese regulatory authorities may interpret it broadly. Furthermore, there remains uncertainty as to whether the Chinese regulatory authorities may adopt new laws, regulations, rules, or detailed implementation and interpretation in relation, or in addition, to the Revised CAC Measures and the Draft Management Regulations. While we intend to closely monitor the evolving laws and regulations in this area and take all reasonable measures to mitigate compliance risks, we cannot guarantee that our business and operations will not be adversely affected by the potential impact of the Revised CAC Measures, the Draft Management Regulations or other laws and regulations related to privacy, data protection and information security. Also, the National People's Congress released the Personal

Information Protection Law, which became effective in 2021. The Personal Information Protection Law provides a comprehensive set of data privacy and protection requirements that apply to the processing of personal information and expands data protection compliance obligations to cover the processing of personal information of persons by organizations and individuals in China, and the processing of personal information of persons in China outside of China if such processing is for purposes of providing products and services to, or analyzing and evaluating the behavior of, persons in China. The Personal Information Protection Law also provides that critical information infrastructure operators and personal information processing entities who process personal information meeting a volume threshold set by Chinese cyberspace regulators are also required to store in China personal information generated or collected in China, and to pass a security assessment administered by Chinese cyberspace regulators for any export of such personal information. Lastly, the Personal Information Protection Law contains proposals for significant fines for serious violations of up to RMB 50 million or 5% of annual revenues from the prior year and may also be ordered to suspend any related activity by competent authorities. We do not maintain, nor do we intend to maintain in the future, personally identifiable health information of patients in China. In addition, certain industry-specific laws and regulations affect the collection and transfer of data in the PRC. The Regulations on the Administration of Human Genetic Resources of the PRC (the “HGR Regulation”), promulgated by the State Council, came into effect in 2019. It stipulates that foreign organizations, individuals, and the entities established or actually controlled by foreign organizations or individuals are forbidden to collect, preserve and export China’s human genetic resources. Foreign organizations and the entities established or actually controlled by foreign organizations or individuals may only utilize and be provided with China’s human genetic resources after satisfaction of all requirements under the HGR Regulation and other applicable laws, such as (i) China’s human genetic resources being utilized only in international cooperation with Chinese scientific research institutions, universities, medical institutions, and enterprises for scientific research and clinical trials after completion of requisite approval or filing formalities with competent governmental authorities, and (ii) China’s human genetic resources information being provided after required filing and information backup procedures have been gone through. In 2020, the SCNPC promulgated the Biosecurity Law of the PRC, which Article 67 reaffirms the regulatory requirements stipulated by the HGR Regulation while potentially increasing the administrative sanctions where China’s human genetic resources are collected, preserved, exported or used in international cooperation in violation of applicable laws. The Ministry of Science and Technology published the Implementing Rules for the Regulations on the Administration of Human Genetic Resources (the “HGR Implementing Rules”), which came into effect in 2023. The HGR Implementing Rules have, among other things, further clarified the scope of China’s human genetic resources information, improved the procedure rules for applicable approval, filing and security review, and refined the provisions with respect to the forbiddance on the collection, preservation and export of China’s human genetic resources by foreign organizations, individuals, and the entities established or actually controlled by foreign organizations or individuals. There remain significant uncertainties as to how various provisions of the HGR Regulation and the related laws and regulations may be interpreted and implemented. Given such uncertainty, although we have made great efforts to comply with mandatory requirements of laws and government authorities in this regard, we cannot assure you that we will be deemed at all times in full compliance with the HGR Regulation, the Biosecurity Law of the PRC, the HGR Implementing Rules and other applicable laws in our utilizing of and dealing with China’s human genetic resources. As a result, we may be exposed to compliance risks under the HGR Regulation, the Biosecurity Law of the PRC and the HGR Implementing Rules. Interpretation, application and enforcement of these laws, rules and regulations evolve from time to time and their scope may continually change, through new legislation, amendments to existing legislation or changes in enforcement. Compliance with China’s new Cyber Security Law and Data Security Law could significantly increase the cost to us of providing our service offerings, require significant changes to our operations or even prevent us from providing certain service offerings in jurisdictions in which we currently operate or in which we may operate in the future. Despite our efforts to comply with applicable laws, regulations and other obligations relating to privacy, data protection and information security, it is possible that our practices, offerings or platform could fail to meet all of the requirements imposed on us by the Cyber Security Law, the Data Security Law and/or related implementing regulations. Any failure on our part to comply with such law or regulations or any other obligations relating to privacy, data protection or information security, or any compromise of security that results in unauthorized access, use or release of personally identifiable information or other data, or the perception or allegation that any of the foregoing types of failure or compromise has occurred, could damage our reputation, discourage new and existing counterparties from contracting with us or result in investigations, fines, suspension or other penalties by Chinese government authorities and private claims or litigation, any of which could adversely affect our business, financial condition and results of operations. Even if our practices are not subject to legal challenge, the perception of privacy concerns, whether or not valid, may harm our reputation and brand and adversely affect our business, financial condition and results of operations. Moreover, the legal uncertainty created by the Data Security Law, the Revised CAC Measures and the recent Chinese government actions could adversely affect our ability, on favorable terms, to raise capital, including engaging in follow-on offerings of our securities in the U.S. market. Pharmaceutical companies operating in China are required to comply with extensive regulations and hold a number of permits and licenses to carry on their business. Our ability to obtain and maintain these regulatory approvals is uncertain, and future government regulation may place additional burdens on our current and planned operations in China. The pharmaceutical industry in China is subject to extensive government regulation and supervision. The regulatory framework addresses all aspects of operating in the pharmaceutical industry, including product development activities, clinical trials, registration, production, distribution, packaging, labeling, storage and shipment, advertising, licensing and post-approval pharmacovigilance certification requirements and procedures, periodic renewal and reassessment processes, data security and data privacy protection requirements and compliance and environmental protection. In particular, we are subject to many of these laws and regulations because our wholly-owned subsidiary, AnHeart Therapeutics (Hangzhou) Co., Ltd., through which we conduct research and development activities, operates primarily in China. Violation of applicable laws and regulations may materially and adversely affect our business. The regulatory framework governing the pharmaceutical industry in China is subject to change and amendment from time to time. Any such change or amendment could materially and adversely impact our business, financial condition and prospects. The Chinese government has introduced various reforms to the Chinese healthcare system in recent years and may continue to do so, with an overall objective to expand basic medical insurance coverage and improve the quality and reliability of healthcare services. The specific regulatory changes under the various reform initiatives remain uncertain. The implementing measures to be issued may not be sufficiently effective to achieve the stated goals, and as a result, we may not be able to benefit from such reform to the extent we expect, if at all. Moreover, the various reform initiatives could give rise to regulatory developments, such as more burdensome administrative procedures, which may have an adverse effect on our business and prospects. As a company with operations and business relationships outside of the United States, our business is subject to economic, political, regulatory and other risks associated with international operations. As a company with operations in China, our business is subject to risks associated with conducting business outside the United States. In addition to our research and development activities through AnHeart Therapeutics (Hangzhou) Co., Ltd. in China, some of our suppliers and clinical trial relationships are located outside the United States. Accordingly, our future results could be harmed by a variety of factors, including: economic weakness, including inflation, or political instability in particular non-U.S. economies and markets; a different and changing regulatory requirements for product approvals; differing jurisdictions could present different issues for securing, maintaining or obtaining freedom to operate in such jurisdictions; potentially reduced protection for intellectual property rights; difficulties in compliance with different, complex and changing laws, regulations and court systems of multiple jurisdictions and compliance with a wide variety of foreign laws, treaties and regulations; changes in non-U.S. regulations and customs, tariffs and trade barriers; changes in non-U.S. currency exchange rates of the RMB; increasing geopolitical tensions between the U.S. and China and changes in a specific country’s or region’s political or economic environment especially with respect to a particular country’s treatment of or stance towards other countries; trade protection measures, import or export licensing requirements or other restrictive actions by governments; differing reimbursement regimes and price controls in certain non-U.S. markets; negative consequences from changes in tax laws; compliance with tax, employment, immigration and labor laws for employees living or traveling abroad; variable tax treatment in different jurisdictions of options granted under our equity incentive plans; workforce uncertainty in countries where labor unrest is more common than in the United States; and business interruptions resulting from geo-political actions, including war and terrorism, health epidemics, or natural disasters including earthquakes, typhoons, floods and fires. If we fail to comply with Chinese environmental, health and safety laws and regulations, we could become subject to fines or penalties or incur costs that could have a material adverse effect on the success of our business. We are subject to numerous environmental, health and safety laws and regulations, including those governing laboratory procedures, fire safety and the handling, use, storage, treatment and disposal of hazardous materials and wastes. Despite our efforts to comply fully with environmental and safety regulations, any violation of these regulations may result in substantial fines, criminal sanctions, revocations of operating permits, the shutdown of our facilities and the incurrance of obligations to take corrective measures. Although we maintain workers’ compensation insurance to cover costs and expenses incurred due to on-the-job injuries to our employees and public liability insurance to cover costs and expenses that may be incurred if third parties are injured on our property, such insurance may not provide adequate coverage against potential liabilities. Furthermore, the Chinese government may take steps towards the adoption of more stringent environmental regulations, and, due to the possibility of unanticipated regulatory or other developments, the amount and timing of future environmental expenditures may vary substantially from those currently anticipated. If there is any unanticipated change in the environmental regulations, our third-party manufacturers and other service providers may incur substantial capital expenditures to install, replace, upgrade or supplement their manufacturing facilities and equipment or make operational changes to limit any adverse impact or potential adverse impact on the environment in order to comply with new environmental protection laws and regulations. If such costs become prohibitively expensive, we may be forced to cease certain aspects of our business operations and our business may be materially adversely affected. Development in the Chinese legal system could materially and adversely affect us. Chinese laws and regulations govern our operations in China and the PRC legal system is a civil law system based on written statutes. Unlike the common law system, prior court decisions under the civil law system may be cited for reference but have limited precedential value. As the laws and regulations are relatively new and the PRC legal system continues to evolve, there may be room for discretion in the implementation of these laws and regulations. And as these laws and regulations are evolving in response to changing economic and other conditions, factors related to the application and implementation of these laws and regulations may affect our business and results of operations. We may be exposed to liabilities under the U.S. Foreign Corrupt Practices Act (the “FCPA”), and similar anti-corruption and anti-bribery laws of China and other countries in which we operate, as well as U.S. and certain foreign export controls, trade sanctions and import laws and regulations. Compliance with these legal requirements could limit our ability to compete in foreign markets and any determination that we have violated these laws could have a material adverse effect on our business or our reputation. Our operations are subject to the FCPA and similar anti-bribery or anti-corruption laws, regulations or rules of China and other countries in which we operate. The FCPA and these other laws generally prohibit us, our officers, and our employees and intermediaries from, directly or indirectly, offering, authorizing or making improper payments to non-U.S. government officials for the purpose of obtaining or retaining business or other advantage. We may engage third parties for clinical trials outside of the United States, to sell our products abroad once we enter a commercialization phase, and/or to obtain necessary permits, licenses, patent registrations and other regulatory approvals. We have direct or indirect interactions with officials and employees of government agencies or government-affiliated hospitals, universities and other organizations. As our business expands, the applicability of the FCPA and other anti-bribery laws to our operations will increase. If our procedures and controls to monitor anti-bribery compliance fail to protect us from reckless or criminal acts committed by our employees or agents or if we, or our employees, agents, contractors or other collaborators, fail to comply with applicable anti-bribery laws, our reputation could be harmed and we could incur criminal or civil penalties, other sanctions and/or significant expenses, which could have a material adverse effect on our business, including our financial condition, results of operations, cash flows and prospects. In addition, our products may be subject to U.S. and foreign export controls, trade sanctions and import laws and regulations. Governmental regulation of the import or export of our products, or our failure to obtain any required import or export authorization for our products, when applicable, could harm our international or domestic sales and adversely affect our revenue. Compliance with applicable regulatory requirements regarding the export of our products may create delays in the introduction of our products in international markets or, in some cases, prevent the export of our products to some countries altogether. Furthermore, U.S. export control laws and economic sanctions prohibit the shipment of certain products and services to countries, governments and persons targeted by U.S. sanctions. If we fail to comply with export and import regulations and such economic sanctions, penalties could be imposed, including fines and/or denial of certain export privileges. Moreover, any new export or import restrictions, new legislation or shifting approaches in the enforcement or scope of existing regulations, or in the countries, persons or products targeted by such regulations, could result in decreased use of our products by, or in our decreased ability to export our products to, existing or potential customers with international operations. Any decreased use of our products or limitation on our ability to export or sell our products would likely adversely affect our business. Regulatory Requirements on currency exchange may limit our ability to receive and use effectively financing in foreign currencies. Our Chinese subsidiaries’ ability to obtain currency exchange is subject to certain foreign exchange regulations and, in the case of transactions under the capital account, requires the approval of and/or registration with Chinese government authorities, including the State Administration of Foreign Exchange (SAFE). In particular, if we finance our Chinese subsidiaries by means of foreign debt from us or other foreign lenders, the amount is not allowed to, among other things, exceed the statutory limits and such loans must be registered with the local branch of SAFE. If we finance our Chinese subsidiaries by means of additional capital contributions, these capital contributions are subject to registration with the State Administration for Market Regulation or its local branch, reporting of foreign investment information with the Ministry of Commerce of the People’s Republic of China (MOFCOM), or its local branch or registration with other governmental authorities in China. In light of the various requirements imposed by Chinese regulations on loans to, and direct investment in, China-based entities by offshore holding companies, we cannot assure you that we will be able to complete the necessary government requirements or obtain the necessary government approvals on a timely basis, if at all, with respect to future loans or capital contributions by us to our Chinese subsidiaries. If we fail to adhere to such requirements or obtain such approval, our ability to fund our Chinese operations, including research and development activities through AnHeart Therapeutics (Hangzhou) Co., Ltd., may be negatively affected, which could materially and adversely affect our ability to fund and expand our business. Chinese regulations relating to the establishment of offshore special purpose companies by residents in China may subject our China resident beneficial owners or our wholly foreign-owned subsidiaries in China to liability or penalties, limit our ability to inject capital into these subsidiaries, limit these subsidiaries’ ability to increase their registered capital or distribute profits to us, or may otherwise adversely affect us. In 2014, SAFE promulgated the SAFE Circular 37, which requires residents of China to register with local branches of SAFE in connection with their direct establishment or indirect control of an offshore entity, for the purpose of overseas investment and financing, with such residents’ legally owned assets or equity interests in domestic enterprises or offshore assets or interests, referred to in SAFE Circular 37 as a “special purpose vehicle.” The term “control” under SAFE Circular 37 is broadly defined as the operation rights, beneficiary rights or decision-making rights acquired by residents of China in the offshore special purpose vehicles or Chinese companies by such means as acquisition, trust, proxy, voting rights, repurchase, convertible bonds or other arrangements. SAFE Circular 37 further requires amendment to the registration in the event of any changes with respect to the basic information of or any significant changes with respect to the special purpose vehicle, such as an increase or decrease of capital contributed by China residents, share transfer or exchange, merger, division or other material events. If the shareholders of the offshore holding company who are residents of China do not complete their registration with the local SAFE branches, the Chinese subsidiaries may be prohibited from making distributions of profits and proceeds from any reduction in capital, share transfer or liquidation to the offshore parent company and from carrying out subsequent cross-border foreign exchange activities, and the offshore parent company may be restricted in its ability to contribute additional capital into its Chinese subsidiaries. Moreover, failure to comply with the SAFE registration and amendment requirements described above could result in liability under Chinese law for evasion of applicable foreign exchange restrictions. Certain residents of China may hold direct or indirect interests in our company, and we will request residents of China who we know hold direct or indirect interests in our company, if any, to make the necessary applications, filings and amendments as required under SAFE Circular 37 and other related rules. However, we may not at all times be fully aware or informed of the identities of our shareholders or beneficial owners that are required to make such registrations, and we cannot provide any assurance that these residents will comply with our requests to make or obtain any applicable registrations or comply with other requirements under SAFE Circular 37 or other related rules. The failure or inability of our China resident shareholders to comply with the registration procedures set forth in these regulations may subject us to fines or legal sanctions, restrictions on our cross-border investment activities or those of our China subsidiaries and

limitations on the ability of our wholly foreign-owned subsidiaries in China to distribute dividends or the proceeds from any reduction in capital, share transfer or liquidation to us, and we may also be prohibited from injecting additional capital into these subsidiaries. Moreover, failure to comply with the various foreign exchange registration requirements described above could result in liability under Chinese law for circumventing applicable foreign exchange restrictions. As a result, our business operations and our ability to make distributions to you could be materially and adversely affected. We and our shareholders face uncertainties in China with respect to indirect transfers of equity interests in China resident enterprises. The indirect transfer of equity interests in China resident enterprises by a non-China resident enterprise (Indirect Transfer), is potentially subject to income tax in China at a rate of 10% on the gain if such transfer is considered as not having a commercial purpose and is carried out for tax avoidance. The State Administration of Taxation (SAT) has issued several rules and notices to tighten the scrutiny over acquisition transactions in recent years. The Announcement of the State Administration of Taxation on Several Issues Concerning the Enterprise Income Tax on Indirect Property Transfer by Non-Resident Enterprises (SAT Circular 7), sets out the scope of Indirect Transfers, which includes any changes in the shareholder's ownership of a foreign enterprise holding Chinese assets directly or indirectly in the course of a group's overseas restructuring, and the factors to be considered in determining whether an Indirect Transfer has a commercial purpose. An Indirect Transfer satisfying all the following criteria will be deemed to lack a bona fide commercial purpose and be taxable under Chinese laws: (i) 75% or more of the equity value of the intermediary enterprise being transferred is derived directly or indirectly from the Chinese taxable assets; (ii) at any time during the one-year period before the indirect transfer, 90% or more of the asset value of the intermediary enterprise (excluding cash) is comprised directly or indirectly of investments in China, or 90% or more of its income is derived directly or indirectly from China; (iii) the functions performed and risks assumed by the intermediary enterprise and any of its subsidiaries that directly or indirectly hold the Chinese taxable assets are limited and are insufficient to prove their economic substance; and (iv) the non-Chinese tax payable on the gain derived from the indirect transfer of the Chinese taxable assets is lower than the potential Chinese income tax on the direct transfer of such assets. A transaction that does not satisfy all four tests in the immediately preceding sentence may nevertheless be deemed to lack a bona fide commercial purpose if the taxpayer cannot justify such purpose from a totality approach, taking into account the transferred group's value, income, asset composition, the history and substance in the structure, the non-Chinese tax implications, any tax treaty benefit and the availability of alternative transactions. Nevertheless, a non-resident enterprise's buying and selling shares of the same listed foreign enterprise on the public market will fall under the safe harbor available under SAT Circular 7 if the shares were purchased on the public market as well and will not be subject to Chinese tax pursuant to SAT Circular 7. We face uncertainties regarding the reporting required for and impact on future private equity financing transactions, share exchanges or other transactions involving the transfer of shares in our company by investors that are non-Chinese resident enterprises, or the sale or purchase of shares in other non-Chinese resident companies or other taxable assets by us. For example, the Chinese tax authorities may consider that a future securities offering involves an indirect change of shareholding in our Chinese subsidiaries and therefore it may be regarded as an Indirect Transfer under SAT Circular 7. Even if we believe no SAT Circular 7 reporting is required on the basis that such an offering has commercial purposes and is not conducted for tax avoidance, Chinese tax authorities may pursue us to report under SAT Circular 7 and request that we and our Chinese subsidiaries assist in the filing. As a result, we and our subsidiaries may be required to expend significant resources to provide assistance and comply with SAT Circular 7, or establish that we or our non-resident enterprises should not be subject to tax under SAT Circular 7, for such an offering or other transactions, which may have an adverse effect on our and their financial condition and day-to-day operations. Any failure to comply with Chinese regulations regarding the registration requirements for our employee equity incentive plans may subject us to fines and other legal or administrative sanctions, which could adversely affect our business, financial condition and results of operations. In 2012, the SAFE promulgated the Notices on Issues Concerning the Foreign Exchange Administration for Domestic Individuals Participating in Stock Incentive Plans of Overseas Publicly Listed Companies (the Stock Option Rules). In accordance with the Stock Option Rules and other relevant rules and regulations, Chinese citizens or non-Chinese citizens residing in China for a continuous period of not less than one year who participate in any stock incentive plan of an overseas publicly listed company, subject to a few exceptions, are required to register with SAFE through a domestic qualified agent, which could be a Chinese subsidiary of such a 71 A overseas listed company, and complete certain procedures. Our employees who are Chinese citizens or who reside in China for a continuous period of not less than one year and who participate in our stock incentive plans are subject to such regulation. We plan to assist our employees to register their equity awards. However, any failure of our Chinese individual beneficial owners and holders of equity awards to comply with the SAFE registration requirements may subject them to fines and legal sanctions and may limit the ability of our Chinese subsidiaries to distribute dividends to us. We also face regulatory uncertainties that could restrict our ability to adopt additional incentive plans for our employees under Chinese law. Risks Related to Ownership of Our Securities The market price of our securities may be volatile and fluctuate substantially, which could result in substantial losses for our investors and may subject us to securities litigation suits. The market price of our securities may be volatile. The stock market in general and the market for pharmaceutical companies in particular have experienced extreme volatility that has often been unrelated to the operating performance of particular companies. As a result of this volatility, investors may not be able to sell their securities or above the price they paid. The market price for our securities may be influenced by many factors, including: adverse regulatory decisions; any delay in our regulatory filings for our product candidates and any adverse development or perceived adverse development with respect to the applicable regulatory authority's review of such filings, including without limitation the FDA's issuance of a refusal to file letter or a request for additional information; the impact of health epidemics; the commencement, enrollment or results of any future clinical trials we may conduct, or changes in the development status of our product candidates; adverse results from, delays in or termination of clinical trials; unanticipated serious safety concerns related to the use of our product candidates; lower than expected market acceptance of our product candidates following approval for commercialization; changes in financial estimates by us or by any securities analysts who might cover our securities; conditions or trends in our industry; changes in the market valuations of similar companies; stock market price and volume fluctuations of comparable companies and, in particular, those that operate in the pharmaceutical industry; publication of research reports about us or our industry or positive or negative recommendations or withdrawal of research coverage by securities analysts; announcements by us or our competitors of significant acquisitions, strategic partnerships or divestitures; announcements of investigations or regulatory scrutiny of our operations or lawsuits filed against us; investors' general perception of our company and our business; recruitment or departure of key personnel; overall performance of the equity markets; trading volume of our securities; currency exchange rate fluctuations; disputes or other developments relating to intellectual property rights, including patents, litigation matters and our ability to obtain, maintain, defend, protect and enforce patent and other intellectual property rights for our technologies; significant lawsuits, including patent or stockholder litigation; proposed changes to healthcare laws in the U.S., China or other foreign jurisdictions, or speculation regarding such changes including changes in the structure of healthcare payment systems; 72 A general political and economic conditions; and other events or factors, many of which are beyond our control. In addition, in the past, stockholders have initiated class action lawsuits against pharmaceutical and biotechnology companies following periods of volatility in the market prices of these companies' stock. Such litigation, if instituted against us, could cause us to incur substantial costs and divert management's attention and resources from our business. The dual-class structure of our Common Stock has the effect of concentrating voting power with our Chief Executive Officer, which limits other stockholders' ability to influence the outcome of important transactions, including a change in control. Dr. Hung holds all of the outstanding shares of our Class B Common Stock and approximately 24% of our Class A and Class B Common Stock outstanding. In addition to voting together with the Class A Common Stock (with one vote per share) on all matters, the holders of Class B Common Stock have (i) the right to elect and remove without cause three of our directors plus at least 50% of all directors in excess of seven and (ii) an approval right over any acquisition (whether by merger, sale of shares or sale of assets) or our liquidation. Accordingly, Dr. Hung has the ability to control or exert substantial influence over all matters submitted to our stockholders for approval, including the election of directors and amendments of our organizational documents, and an approval right over any acquisition or liquidation of our company. Dr. Hung may have interests that differ from those of the other stockholders and may vote in a way with which the other stockholders disagree and which may be adverse to their interests. This concentrated control may have the effect of delaying, preventing or deterring a change in control, could deprive our stockholders of an opportunity to receive a premium for their capital stock as part of a sale of our company, and might ultimately affect the market price of shares of our Class A Common Stock. We cannot predict the impact our dual-class structure may have on the market price of our Class A Common Stock. We cannot predict whether our dual-class structure, combined with the concentrated voting power of Dr. Hung by virtue of his ownership of 100% of the outstanding shares of our Class B Common Stock, will result in a lower or more volatile market price of our Class A Common Stock in the future, or in adverse publicity or other adverse consequences. Certain index providers have announced restrictions on including companies with multi-class share structures in certain of their indices. For example, in 2017, FTSE Russell and Standard & Poor's announced that they would cease to allow most newly public companies utilizing dual or multi-class capital structures to be included in their indices. Under the announced policies, our dual-class capital structure makes us ineligible for inclusion in any of these indices. Given the sustained flow of investment funds into passive strategies that seek to track certain indices, exclusion from stock indices would likely preclude investment by many of these funds and could make our securities less attractive to other investors. As a result, the market price of our Class A Common Stock could be adversely affected. There can be no assurance that we will be able to comply with the continued listing standards of the NYSE. Our Class A Common Stock and warrants are listed on the NYSE under the symbols NUVB and NUVBW, respectively. Our continued eligibility for listing will depend on our compliance with the continued listing standards of the NYSE and may depend on the number of our shares that are redeemed. If the NYSE delists our securities from trading on its exchange for failure to meet the listing standards, we and our stockholders could face significant negative consequences including: limited availability of market quotations for our securities; a determination that our Common Stock is a penny stock which will require brokers trading in our Common Stock to adhere to more stringent rules, possibly resulting in a reduced level of trading activity in the secondary trading market for shares of our Common Stock; a limited amount of analyst coverage; and a decreased ability to issue additional securities or obtain additional financing in the future. Future sales, or the perception of future sales, by us or our stockholders in the public market could cause the market price for our securities to decline. The sale of our securities in the public market, or the perception that such sales could occur, could harm the prevailing market price of our securities. These sales, or the possibility that these sales may occur, also might make it more difficult for us to sell equity securities in the future at a time and at a price that we deem appropriate. Immediately prior to our Acquisition of AnHeart, we had a total of approximately 219,083,219 shares of Common Stock outstanding, consisting of approximately 218,083,219 shares of Class A Common Stock and 1,000,000 shares of Class B Common Stock. All of these shares are freely tradable without registration under the Securities Act, and without restriction by persons other than our affiliates (as defined under Rule 144 of the Securities Act, Rule 144), including our directors, executive officers and other affiliates. 73 A In the AnHeart Acquisition, we issued to AnHeart securityholders approximately (i) 27,646,255 shares of Class A Common Stock, (ii) shares of preferred stock convertible into an aggregate of approximately 85,120,200 shares of Class A Common Stock upon the approval of such conversion by our stockholders, and warrants exercisable for approximately 2,893,731 shares of Class A Common Stock at an exercise price of \$11.50 per share. These securities were issued in a private placement and are subject to restrictions on resale under federal securities laws and, in most cases, lockup agreements restricting resale of more than 20% of such securities before December 31, 2024. We also reserved an aggregate of approximately 15,943,933 shares of Class A Common Stock for issuance under AnHeart's equity incentive plans. We intend to file with the SEC a registration statement with respect to the resale of the shares and warrants issued in the AnHeart Acquisition. In addition, the shares of Class A Common Stock reserved for future issuance under our equity incentive plans will become eligible for sale in the public market once those shares are issued, subject to provisions relating to various vesting agreements, lock-up agreements and, in some cases, limitations on volume and manner of sale applicable to affiliates under Rule 144, as applicable. Our compensation committee of our board of directors may determine the exact number of shares to be reserved for future issuance under our equity incentive plans at its discretion. We have filed and expect to file registration statements on Form S-8 under the Securities Act to register shares of Class A Common Stock or securities convertible into or exchangeable for shares of Class A Common Stock issued pursuant to our equity incentive plans. Any such Form S-8 registration statements will automatically become effective upon filing. Accordingly, shares registered under such registration statements will be available for sale in the open market. In the future, we may also issue our securities in connection with investments or acquisitions. The amount of shares of Class A Common Stock issued in connection with an investment or acquisition could constitute a material portion of our then-outstanding shares of Class A Common Stock. Any issuance of additional securities in connection with investments or acquisitions may result in additional dilution to our stockholders. Because we do not anticipate paying any cash dividends on our Class A Common Stock in the foreseeable future, capital appreciation, if any, will be your sole source of gains and you may never receive a return on your investment. We may retain future earnings, if any, for future operations, expansion and debt repayment and have no current plans to pay any cash dividends for the foreseeable future. Any decision to declare and pay dividends as a public company in the future will be made at the discretion of our board of directors and will depend on, among other things, our results of operations, financial condition, cash requirements, contractual restrictions and other factors that our board of directors may deem relevant. In addition, our ability to pay dividends may be limited by covenants of any existing and future outstanding indebtedness we or our subsidiaries incur. As a result, you may not receive any return on an investment in our securities unless you sell your securities for a price greater than that which you paid for it. There is no guarantee that our warrants will be in the money at the time they become exercisable, and they may expire worthless. The exercise price for our outstanding warrants is \$11.50 per share of Class A Common Stock. There is no guarantee that any of our warrants will be in the money following the time they become exercisable and prior to their expiration, and as such, the warrants may expire worthless. We may issue additional securities without your approval, which would dilute your ownership interests and may depress the market price of our securities. As of June 30, 2024, we have options outstanding to purchase approximately 52,692,948 shares of Class A Common Stock. Pursuant to the 2021 Equity Incentive Plan (the 2021 Plan) and the Employee Stock Purchase Plan (the 2021 ESPP), we may issue under the 2021 Plan an aggregate of up to 54,911,238 shares of Class A Common Stock and Class B Common Stock, which amount will be subject to increase from time to time. In addition, in the AnHeart Acquisition, we assumed the AnHeart equity incentive plans and reserved an aggregate of approximately 15,943,933 shares of Class A Common Stock for issuance upon exercise of outstanding options or settlement of outstanding restricted stock units issued under those plans. We may also issue additional shares of Class A Common Stock or other equity securities of equal or senior rank in the future in connection with, among other things, future acquisitions or repayment of outstanding indebtedness, without stockholder approval, in a number of circumstances. The issuance of additional shares or other equity securities of equal or senior rank would have the following effects: existing stockholders' proportionate ownership interest in our company will decrease; the amount of cash available per share, including for payment of dividends in the future, may decrease; the relative voting strength of each previously outstanding Common Stock may be diminished; and the market price of our securities may decline. 74 A Anti-takeover provisions in our amended and restated certificate of incorporation and under Delaware law could make an acquisition of our company, which may be beneficial to our stockholders, more difficult, and may prevent attempts by our stockholders to replace or remove our current management. Our amended and restated certificate of incorporation contains provisions that may delay or prevent an acquisition of the company or change in our management in addition to the significant rights of Dr. Hung as the holder of 100% of the outstanding shares of our Class B Common Stock. These provisions may make it more difficult for stockholders to replace or remove members of our board of directors. Because the board of directors is responsible for appointing the members of the management team, these provisions could in turn frustrate or prevent any attempt by our stockholders to replace or remove our current management. In addition, these provisions could limit the price that investors might be willing to pay in the future for shares of our Class A Common Stock. Among other things, these provisions include: the limitation of the liability of, and the indemnification of, our directors and officers; a prohibition on actions by our stockholders except at an annual or special meeting of stockholders; a prohibition on actions by our stockholders by written consent; and the ability of the board of directors to issue preferred stock without stockholder approval, which could be used to institute a poison pill that would work to dilute the stock ownership of a potential hostile acquirer, effectively preventing acquisitions that have not been approved by the board of directors. Moreover, because we are incorporated in Delaware, we are governed by the provisions of Section 203 of the Delaware General Corporation Law (the DGCL), which prohibits a person who owns 15% or more of our outstanding voting stock from merging or combining with us for a period of three years after the date of the transaction in which the person acquired 15% or more of our outstanding voting stock, unless the merger or combination is approved in a prescribed manner. This could discourage, delay or prevent a third party from acquiring or merging with us, whether or not it is desired by, or beneficial to, our stockholders. This

could also have the effect of discouraging others from making tender offers for our Class A common stock, including transactions that may be in our stockholders' best interests. Finally, these provisions establish advance notice requirements for nominations for election to the board of directors or for proposing matters that can be acted upon at stockholder meetings. These provisions would apply even if the offer may be considered beneficial by some stockholders. Our amended and restated certificate of incorporation provides that the Court of Chancery of the State of Delaware and the federal district courts of the United States of America are the exclusive forums for substantially all disputes between us and our stockholders, which could limit our stockholders' ability to obtain a favorable judicial forum for disputes with us or our directors, officers or employees. Our amended and restated certificate of incorporation provides that the Court of Chancery of the State of Delaware will be the exclusive forum for the following types of actions or proceedings under Delaware statutory or common law: any derivative action or proceeding brought on our behalf; any action asserting a breach of fiduciary duty; any action asserting a claim against us arising under the Delaware General Corporation Law, our amended and restated certificate of incorporation or our amended and restated bylaws; and any action asserting a claim against us that is governed by the internal-affairs doctrine or otherwise related to our internal affairs. To prevent having to litigate claims in multiple jurisdictions and the threat of inconsistent or contrary rulings by different courts, among other considerations, our amended and restated certificate of incorporation further provides that the federal district courts of the United States of America will be the exclusive forum for resolving any complaint asserting a cause of action arising under the Securities Act. This provision would not apply to suits brought to enforce a duty or liability created by the Exchange Act. Furthermore, Section 22 of the Securities Act creates concurrent jurisdiction for federal and state courts over all such Securities Act actions. Accordingly, both state and federal courts have jurisdiction to entertain such claims. While the Delaware courts have determined that such choice of forum provisions are facially valid, a stockholder may nevertheless seek to bring a claim in a venue other than those designated in the exclusive forum provisions. In such instance, we would expect to vigorously assert the validity and enforceability of the exclusive forum provisions of our amended and restated certificate of incorporation. This may require significant additional costs associated with resolving such action in other jurisdictions and there can be no assurance that the provisions will be enforced by a court in those other jurisdictions. These exclusive forum provisions may limit a stockholder's ability to bring a claim in a judicial forum that it finds favorable for disputes with us or our directors, officers or other employees, which may discourage lawsuits against us and our directors, officers and other employees. If a court were to find either exclusive-forum provision in our amended and restated certificate of incorporation to 75 Å to be inapplicable or unenforceable in an action, we may incur further significant additional costs associated with resolving the dispute in other jurisdictions, all of which could harm our business. We are eligible to report as a smaller reporting company, and as a result of the reduced reporting requirements applicable to smaller reporting companies, our securities may be less attractive to investors. We are eligible to report as a smaller reporting company. For as long as we continue to be eligible to report as a smaller reporting company, we may take advantage of exemptions from various reporting requirements that are applicable to other public companies that are not smaller reporting companies, including exemptions from the requirements of holding a nonbinding advisory vote on executive compensation and stockholder approval of any golden parachute payments not previously approved. If some investors find our securities less attractive because we rely on any of these exemptions, there may be a less active trading market for our securities and the price of our securities may be more volatile. General Risk Factors If we fail to maintain proper and effective internal controls, our ability to produce accurate financial statements on a timely basis could be impaired. We are subject to the reporting requirements of the Securities Exchange Act of 1934, as amended, or the Exchange Act, the Sarbanes-Oxley Act and the rules and regulations of the NYSE. Section 302 of the Sarbanes-Oxley Act requires, among other things, that public companies report on the effectiveness of our disclosure controls and procedures in our quarterly and annual reports and, beginning with this report, Section 404 of the Sarbanes-Oxley Act requires that we perform system and process evaluation and testing of our internal control over financial reporting to allow management to report on the effectiveness of our internal control over financial reporting in our Annual Report on Form 10-K for that year. This has required us to incur substantial additional professional fees and internal costs to expand our accounting and finance functions and to expend significant management efforts. If we are not able to comply with the requirements of Section 404 of the Sarbanes-Oxley Act in a timely manner, or if we are unable to maintain proper and effective internal controls, we may not be able to produce timely and accurate financial statements. If that were to happen, the market price of our stock could decline and we could be subject to sanctions or investigations by the NYSE, the SEC or other regulatory authorities. In addition, our securities may not be able to remain listed on the NYSE or any other securities exchange. We will incur costs and demands upon our management as a result of complying with the laws and regulations affecting public companies in the U.S., which may harm our business. As a public company listed in the U.S., we incur on an ongoing basis significant legal, accounting and other expenses. In addition, changing laws, regulations and standards relating to corporate governance and public disclosure, including regulations implemented by the SEC and the NYSE may increase legal and financial compliance costs and make some activities more time consuming. These laws, regulations and standards are subject to varying interpretations and, as a result, their application in practice may evolve over time as new guidance is provided by regulatory and governing bodies. We intend to invest resources to comply with evolving laws, regulations and standards, and this investment may result in increased general and administrative expenses and a diversion of management's time and attention from regular business activities to compliance activities. If, notwithstanding our efforts, we fail to comply with new laws, regulations and standards, regulatory authorities may initiate legal proceedings against us and our business may be harmed. Failure to comply with these rules might also make it more difficult for us to obtain certain types of insurance, including director and officer liability insurance, and we might be forced to accept reduced policy limits and coverage or incur substantially higher costs to obtain the same or similar coverage. The impact of these events could also make it more difficult for us to attract and retain qualified persons to serve on our board of directors, on committees of our board of directors or as members of senior management. If securities or industry analysts cease publishing research or reports about us, our business or our market, or if they change their recommendations regarding our securities adversely, the price and trading volume of our securities could decline. Equity research analysts may cease providing research coverage of our securities at any time, and such lack of research coverage may adversely affect the market price of our securities. In any event, we do not have any control over the analysts or the content and opinions included in their reports and the price of our stock could decline if one or more equity research analysts downgrade our stock or issue other unfavorable commentary or research. If one or more equity research analysts ceases coverage of our company or fails to publish reports on us regularly, demand for our stock could decrease, which in turn could cause our securities' prices or trading volume to decline. Å 76 Å Item 2. Unregistered Sales of Equity Securities and Use of Proceeds. (a) Issuer Purchases of Equity Securities Å Period Å Total number of shares (or units) purchased Å Average price paid per share (or unit) Å Total number of shares (or units) purchased as part of publicly announced plans or programs Å Maximum number (or approximated dollar value) of shares (or units) that may yet be purchased under the plans or programs April 2024 Å N/A Å \$ N/A Å N/A May 2024 Å N/A Å \$ N/A Å N/A June 2024 Å N/A Å \$ N/A Å N/A July 2024 Å N/A Å \$ N/A Å N/A August 2024 Å N/A Å \$ N/A Å N/A September 2024 Å N/A Å \$ N/A Å N/A October 2024 Å N/A Å \$ N/A Å N/A November 2024 Å N/A Å \$ N/A Å N/A December 2024 Å N/A Å \$ N/A Å N/A Item 3. Defaults Upon Senior Securities. Not applicable. Item 4. Mine Safety Disclosures. Not applicable. Item 5. Other Information Including Trading Arrangements On June 21, 2024, Jerry Wang, AnHeart's Chief Executive Officer and member of the board of directors, entered into a trading plan intended to satisfy the affirmative defense conditions of Rule 10b5-1(c) under the Exchange Act, providing for the sale of up to 130,886 shares of the Company's Common Stock. Pursuant to this plan, Mr. Wang may sell shares beginning October 21, 2024 and ending November 1, 2025. On June 21, 2024, Kerry Wentworth, the Company's Chief Regulatory Officer, entered into a trading plan intended to satisfy the affirmative defense conditions of Rule 10b5-1(c) under the Exchange Act, providing for the sale of up to 303,920 shares of the Company's Common Stock. Pursuant to this plan, Ms. Wentworth may sell shares beginning October 21, 2024 and ending November 1, 2025. On June 21, 2024, David Hanley, the Company's Chief Technical Operations Officer, entered into a trading plan intended to satisfy the affirmative defense conditions of Rule 10b5-1(c) under the Exchange Act, providing for the sale of up to 500,000 shares of the Company's Common Stock. Pursuant to this plan, Mr. Hanley may sell shares beginning October 21, 2024 and ending November 1, 2025. On June 21, 2024, Gary Hattersley, the Company's Chief Scientific Officer, entered into a trading plan intended to satisfy the affirmative defense conditions of Rule 10b5-1(c) under the Exchange Act, providing for the sale of up to 398,935 shares of the Company's Common Stock. Pursuant to this plan, Mr. Hanley may sell shares beginning October 21, 2024 and ending November 1, 2026. Å Item 6. Exhibits. The following exhibits are filed as part of this report: Å Exhibit Number Description 2.1 Å Agreement and Plan of Merger and Reorganization, dated March 24, 2024, by and among Nuvation Bio Inc., AnHeart Therapeutics Ltd., Artemis Merger Sub I, Ltd. and Artemis Merger Sub II, Ltd. (incorporated by reference to Exhibit 2.1 to the Company's Current Report on Form 8-K (File No. 001-39351) filed on March 25, 2024) Å 77 Å Å 3.1 Å Amended and Restated Certificate of Incorporation previously filed with the SEC on February 12, 2021 as Exhibit 3.1 to the Company's Current Report on Form 8-K (File No. 001-39351), which is incorporated herein by reference. Å Å 3.2 Å Amended and Restated Bylaws previously filed with the SEC on February 12, 2021 as Exhibit 3.2 to the Company's Current Report on Form 8-K (File No. 001-39351), which is incorporated herein by reference. Å Å 4.1 Å Form of Certificate of Designation of Preferences, Rights and Limitations of Series A Non-Voting Convertible Preferred Stock previously filed with the SEC on March 25, 2024 as Exhibit 3.1 to the Company's Current Report on Form 8-K (File No. 001-39351), which is incorporated herein by reference. 10.1# Å Non-Employee Director Compensation Policy previously filed with the SEC on May 14, 2024 as Exhibit 10.1 to the Company's Quarterly Report on Form 10-Q (File No. 001-39351), which is incorporated herein by reference. Å Å 10.2# Å Forms of Option Grant Notice and Option Agreement under the 2021 Equity Incentive Plan Å Long-Term Incentive Program previously filed with the SEC on May 9, 2022 as Exhibit 10.2 to the Company's Quarterly Report on Form 10-Q (File No. 001-39351), which is incorporated herein by reference. Å Å 10.3Å; Å License Agreement between Daiichi Sankyo Company, Limited and AnHeart Therapeutics Inc., dated December 7, 2018 previously filed with the SEC on May 14, 2024 as Exhibit 10.3 to the Company's Quarterly Report on Form 10-Q (File No. 001-39351), which is incorporated herein by reference. Å Å 10.4Å; Å First Amendment to License Agreement between Daiichi Sankyo Company, Limited and AnHeart Therapeutics Inc., dated August 17, 2024 previously filed with the SEC on May 14, 2024 as Exhibit 10.4 to the Company's Quarterly Report on Form 10-Q (File No. 001-39351), which is incorporated herein by reference. Å Å 10.5Å; Å License Agreement between Daiichi Sankyo Company, Limited and AnHeart Therapeutics Inc., dated September 7, 2020 previously filed with the SEC on May 14, 2024 as Exhibit 10.5 to the Company's Quarterly Report on Form 10-Q (File No. 001-39351), which is incorporated herein by reference. Å Å 31.1* Certification of Principal Executive Officer Pursuant to Rules 13a-14(a) and 15d-14(a) under the Securities Exchange Act of 1934, as Adopted Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002. Å Å 31.2* Certification of Principal Financial Officer Pursuant to Rules 13a-14(a) and 15d-14(a) under the Securities Exchange Act of 1934, as Adopted Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002. Å Å 32.1* Certification of Principal Executive Officer Pursuant to 18 U.S.C. Section 1350, as Adopted Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002. Å Å 101.INS Inline XBRL Instance Document 101.SCH Å Inline XBRL Taxonomy Extension Schema Document 104 Å Cover Page Interactive Data File (embedded within the Inline XBRL document) Å # Indicates management contract or compensatory plan or arrangement. Å Certain portions of this exhibit will be omitted because they are not material and would likely cause competitive harm to the registrant if disclosed. Å Confidential portions of this Exhibit were redacted pursuant to Item 601(b)(10) of Regulation S-K and the Company agrees to furnish supplementally to the Securities and Exchange Commission a copy of any omitted schedule and/or exhibit upon request. * Filed herewith. Å 78 Å SIGNATURES Pursuant to the requirements of the Securities Act of 1933, the registrant has duly caused this registration statement to be signed on its behalf by the undersigned, thereunto duly authorized in the City of New York, State of New York, on this 5th day of August, 2024. Å Å NUVATION BIO INC. Å Å Date: August 5, 2024 /s/ David Hung, M.D. Å David Hung, M.D. Å Founder, President and Chief Executive Officer Å Å Date: August 5, 2024 /s/ Moses Makunje Å Moses Makunje, VP, Finance Principal Financial and Accounting Officer Å 79 EX-31.1 2 nuvb-ex31.1.htm EX-31.1 EX-31.1 EXHIBIT 31.1 CERTIFICATION PURSUANT TO RULES 13a-14(a) AND 15d-14(a) UNDER THE SECURITIES EXCHANGE ACT OF 1934, AS ADOPTED PURSUANT TO SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002 I, David Hung, M.D. certify that: 1.I have reviewed this Quarterly Report on Form 10-Q of Nuvation Bio, Inc.; 2.Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report; 3.Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report; 4.The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have: (a)Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared; (b)Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles; (c)Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and (d)Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and 5.The registrant's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions): (a)All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and (b)Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting. Å Date: August 5, 2024 By: /s/ David Hung, M.D. David Hung, M.D. Founder, President and Chief Executive Officer (Principal Executive Officer) Å EX-31.2 3 nuvb-ex31.2.htm EX-31.2 EX-31.2 EXHIBIT 31.2 CERTIFICATION PURSUANT TO RULES 13a-14(a) AND 15d-14(a) UNDER THE SECURITIES EXCHANGE ACT OF 1934, AS ADOPTED PURSUANT TO SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002 I, Moses Makunje, certify that: 1.I have reviewed this Quarterly Report on Form 10-Q of Nuvation Bio, Inc.; 2.Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report; 3.Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report; 4.The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have: (a)Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared; (b)Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles; (c)Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and (d)Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and 5.The registrant's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions): (a)All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and (b)Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's

internal control over financial reporting. Â Date: August 5, 2024 By: /s/ Moses Makunje Moses MakunjeVP, Finance(Principal Financial and Accounting Officer) Â Â Â EX-32.1 4 nuvb-ex32.1.htm EX-32.1 EX-32.1 Â Exhibit 32.1 CERTIFICATION PURSUANT TO 18 U.S.C. SECTION 1350, AS ADOPTED PURSUANT TO SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002 In connection with the Quarterly Report of Nuvation Bio, Inc. (the âCompanyâ) on Form 10-Q for the period ending June 30, 2024 as filed with the Securities and Exchange Commission on the date hereof (the âReportâ), I certify, pursuant to 18 U.S.C. Â§ 1350, as adopted pursuant to Â§ 906 of the Sarbanes-Oxley Act of 2002, that: (1) The Report fully complies with the requirements of section 13(a) or 15(d) of the Securities Exchange Act of 1934; and (2) The information contained in the Report fairly presents, in all material respects, the financial condition and result of operations of the Company. Â Date: August 5, 2024 By: /s/ David Hung, M.D. David Hung, M.D. Founder, President and Chief Executive Officer (Principal Executive Officer) Â Â EX-32.2 5 nuvb-ex32_2.htm EX-32.2 EX-32.2 Â Exhibit 32.2 CERTIFICATION PURSUANT TO 18 U.S.C. SECTION 1350, AS ADOPTED PURSUANT TO SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002 In connection with the Quarterly Report of Nuvation Bio, Inc. (the âCompanyâ), on Form 10-Q for the period ending June 30, 2024 as filed with the Securities and Exchange Commission on the date hereof (the âReportâ), I certify, pursuant to 18 U.S.C. Â§ 1350, as adopted pursuant to Â§ 906 of the Sarbanes-Oxley Act of 2002, that: (1) The Report fully complies with the requirements of section 13(a) or 15(d) of the Securities Exchange Act of 1934; and (2) The information contained in the Report fairly presents, in all material respects, the financial condition and result of operations of the Company. Â Date: August 5, 2024 By: /s/ Moses Makunje Moses MakunjeVP, Finance(Principal Financial and Accounting Officer) Â Â EX-101.SCH 6 nuvb-20240630.xsd XBRL TAXONOMY EXTENSION SCHEMA WITH EMBEDDED LINKBASES DOCUMENT 100000 - Document - Document and Entity Information link.presentationLink link:calculationLink link:definitionLink link:definitionLink 100010 - Statement - Consolidated Balance Sheets (Unaudited) link.presentationLink link:calculationLink link:definitionLink 100020 - Disclosure - Leases - Schedule of Aggregate Lease Payments (Details) 2 link.presentationLink link:calculationLink link:definitionLink 100030 - Statement - Consolidated Balance Sheets (Parenthetical) (Unaudited) link.presentationLink link:calculationLink link:definitionLink 100040 - Statement - Consolidated Statements of Cash Flows (Unaudited) 2 link.presentationLink link:calculationLink link:definitionLink 100050 - Statement - Consolidated Statements of Operations and Comprehensive Loss (Unaudited) 2 link.presentationLink link:calculationLink link:definitionLink 100060 - Statement - Consolidated Statements of Operations and Comprehensive Loss (Unaudited) link.presentationLink link:calculationLink link:definitionLink 100070 - Statement - Consolidated Statements of Redeemable Convertible Preferred Stock and Stockholders' Equity (Unaudited) link.presentationLink link:calculationLink link:definitionLink 100080 - Statement - Consolidated Statements of Cash Flows (Unaudited) link.presentationLink link:calculationLink link:definitionLink 995455 - Disclosure - Nature of Operations link.presentationLink link:calculationLink link:definitionLink 995465 - Disclosure - Significant Accounting Policies link.presentationLink link:calculationLink link:definitionLink 995475 - Disclosure - Acquisition link.presentationLink link:calculationLink link:definitionLink 995485 - Disclosure - Fair Value Measurements and Marketable Securities Available-for-Sale link.presentationLink link:calculationLink link:definitionLink 995495 - Disclosure - Collaboration and License Agreements link.presentationLink link:calculationLink link:definitionLink 995505 - Disclosure - Borrowings link.presentationLink link:calculationLink link:definitionLink 995515 - Disclosure - Stockholders' Equity link.presentationLink link:calculationLink link:definitionLink 995525 - Disclosure - Leases link.presentationLink link:calculationLink link:definitionLink 995535 - Disclosure - Net Loss Per Share link.presentationLink link:calculationLink link:definitionLink 995545 - Disclosure - Accumulated Other Comprehensive Income link.presentationLink link:calculationLink link:definitionLink 995555 - Disclosure - Stock-Based Compensation link.presentationLink link:calculationLink link:definitionLink 995565 - Disclosure - Warrants link.presentationLink link:calculationLink link:definitionLink 995575 - Disclosure - Commitments and Contingencies link.presentationLink link:calculationLink link:definitionLink 995585 - Disclosure - Subsequent Event link.presentationLink link:calculationLink link:definitionLink 995595 - Disclosure - Significant Accounting Policies (Policies) link.presentationLink link:calculationLink link:definitionLink 995605 - 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Warrants (Tables) link.presentationLink link:calculationLink link:definitionLink 995695 - Disclosure - Nature of Operations - Additional Information (Details) link.presentationLink link:calculationLink link:definitionLink 995705 - Disclosure - Significant Accounting Policies - Additional Information (Details) link.presentationLink link:calculationLink link:definitionLink 995715 - Disclosure - Significant Accounting Policies - Schedule of Concentration of Credit Risk Net Revenue (Details) link.presentationLink link:calculationLink link:definitionLink 995725 - Disclosure - Significant Accounting Policies - Schedule of Concentration of Customers and Suppliers Accounts Receivable (Details) link.presentationLink link:calculationLink link:definitionLink 995735 - Disclosure - Significant Accounting Policies - Schedule of Significant Accounting Policies Concentration of Risk Suppliers Represent Total Purchases (Details) link.presentationLink link:calculationLink link:definitionLink 995745 - Disclosure - 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Disclosure - Collaboration and License Agreements - Schedule Of Contract Assets And Contract Liabilities (Details) link.presentationLink link:calculationLink link:definitionLink 995865 - Disclosure - Borrowings (Additional Information) (Details) link.presentationLink link:calculationLink link:definitionLink 995875 - Disclosure - Leases - Schedule of Aggregate Lease Payments (Details) link.presentationLink link:calculationLink link:definitionLink 995885 - Disclosure - Leases - (Additional Information) (Details) link.presentationLink link:calculationLink link:definitionLink 995895 - Disclosure - Net Loss Per Share - Summary of Securities Outstanding Excluded from Calculation of Weighted Average Shares Outstanding (Details) link.presentationLink link:calculationLink link:definitionLink 995905 - Disclosure - Accumulated Other Comprehensive Income (Loss) - Schedule of Unrealized Gains (Losses) on available-for-sale securities (Details) link.presentationLink link:calculationLink link:definitionLink 995915 - 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Disclosure - Warrants - Summary of Changes in Liabilities Classified in Level 3 of the Fair Value Hierarchy (Details) link.presentationLink link:calculationLink link:definitionLink 995995 - Disclosure - Commitments and Contingencies - Additional Information (Details) link.presentationLink link:calculationLink link:definitionLink 996005 - Disclosure - Subsequent Event (Additional Information) (Details) link.presentationLink link:calculationLink link:definitionLink Collaborative Arrangement Collaborative Arrangement, Accounting Policy [Policy Text Block] Statement [Line Items] Statement [Line Items] Debt Securities, Held-to-Maturity, Allowance for Credit Loss [Line Items] Accounts receivable Increase (Decrease) in Accounts Receivable Plan Name Plan Name [Domain] Total financial assets: Total financial assets: Total financial assets New York City. New York City [Member] Supplier A [Member] Supplier A [Member] Supplier A [Member] Schedule of Stockholders' Equity Note, Warrants or Rights [Table Text Block] Schedule of Fair Values for Outstanding Warrants Entity Address, Postal Zip Code Entity Address, Postal Zip Code Lessee Operating Lease Expiration Year Lessee operating lease expiration year. Lessee Operating Lease Expiration Year Unrealized gain (loss) Marketable Security, Unrealized Gain (Loss) Marketable Securities, Unrealized Gain (Loss), Total Entity Incorporation, State or Country Code Entity Incorporation, State or Country Code Common Stock Including Additional Paid in Capital [Member] Common Stock Including Additional Paid In Capital [Member] Acquisition Business Combination Disclosure [Text Block] Research and development asset Research and Development Asset Acquired in Transaction Other than Business Combination or Joint Venture Formation, Capitalized Cost Fair Value Equity Level1 To Level2 or Level3 Transfers Description Fair Value Equity Level1 To Level2 or Level3 Transfers Description Shares, Outstanding Balance (in Shares) Balance (in Shares) Fair Value Hierarchy and NAV Fair Value Hierarchy and NAV [Axis] Nonrecurring Adjustment [Axis] Level 2 [Member] Fair Value, Inputs, Level 2 [Member] Share Based Compensation Arrangement By Share Based Payment Award [Line Items] Share-Based Compensation Arrangement by Share-Based Payment Award [Line Items] Subordinated Borrowing [Line Items] Area of Real Estate Property Area of Office Space Total Mezzanine Equity, Total Total mezzanine equity Total Mezzanine Equity Total Mezzanine Equity Lessee, Operating Leases Leases Commitments and Contingencies Commitments and Contingencies Disclosure [Text Block] Derivative Liability, Fair Value, Gross Liability, Total Derivative Liability, Fair Value, Gross Liability Warrants Marketable Security, Gain (Loss) [Abstract] Net realized gains (losses) on available-for-sale securities were as follows: Product and Service [Axis] Collaborative Arrangement and Arrangement Other than Collaborative [Domain] Document Information [Table] Concentration Risk Benchmark [Axis] Stockholders' Equity Equity [Text Block] Class of Warrant or Right Class of Warrant or Right [Axis] Cash and Cash Equivalents [Domain] Debt Securities, Available-for-Sale, Unrealized Loss Position, Fair Value [Table Text Block] Schedule of Marketable Securities with Continuous Unrealized Losses Entity Central Index Key Entity Central Index Key Cash Equivalents, at Carrying Value, Total Cash Equivalents, at Carrying Value Cash equivalents Market Risk Benefit, Change in Fair Value, Gain (Loss) Change in Fair Value, Gain (Loss) Change in Fair Value, Gain (Loss) Warrant [Member] Warrants [Member] Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Forfeitures in Period Number of share option, Forfeited Forfeited Increase (Decrease) in Operating Capital [Abstract] Change in operating assets and liabilities (net of assets and liabilities acquired in the Merger) Concentration Risk [Table] Common stock and additional paid in capital, shares issued Common Stock, Shares, Issued Common Stock, Shares, Issued, Total Share issued fully paid up Marketable securities [Member] Marketable securities [Member] Marketable securities [Member] Schedule Of Employee Service Share Based Compensation Allocation Of Recognized Period Costs [Table] Share-Based Payment Arrangement, Expensed and Capitalized, Amount [Table] China and Japan [Member] China and Japan [Member] China and Japan [Member] San Francisco. San Francisco [Member] Income Statement [Abstract] Business Acquisition, Name of Acquired Entity Entity Address, State or Province Entity Address, State or Province Loss before income taxes Income (Loss) from Continuing Operations before Income Taxes, Noncontrolling Interest Class Of Warrant Or Right [Table] Class of Warrant or Right [Table] Warrants issued Stock and Warrants Issued During Period Share Preferred Stock and Warrants Stock and Warrants Issued During Period Share Preferred Stock and Warrants Issuance of convertible preferred stock related to the AnHeart acquisition Issuance of Convertible Preferred Stock related to the AnHeart Acquisition Issuance of Convertible Preferred Stock related to the AnHeart Acquisition Capitalized contract cost Capitalized Contract Cost, Net Capitalized Contract Cost, Net, Total Loan and Security Agreement [Member] Loan and Security Agreement [Member] Loan and Security Agreement [Member] Equity Securities, FV-NI, Unrealized Loss Net realized losses on equities sold Class A Common Shares Subject to Repurchase. Class A Common Shares Subject to Repurchase [Member] Concentration Risk Type [Domain] Loss Contingencies [Table] Loss Contingencies [Table] Short-term borrowings Short-Term Debt Short-Term Debt, Total Marketable securities Beginning Balance Ending Balance Marketable Securities Marketable Securities, Total Contract with customer liability revenue recognized Contract liabilities Contract with Customer, Liability, Revenue Recognized Customer Concentration Risk [Member] Customer Concentration Risk [Member] Purchases of property and equipment Payments to Acquire Machinery and Equipment Debt Securities, Available-for-sale, Unrealized Loss Position, Total Debt Securities, Available-for-Sale, Unrealized Loss Position Debt Securities, Available-for-sale, Unrealized Loss Position Schedule of Realized Gain (Loss) [Table Text Block] Schedule of Net Realized Gain (Loss) on Marketable Securities Preferred Stock, Convertible, Shares Issuable Number of convertible preferred stock converted to common stock Interest receivable on marketable securities Increase (Decrease) in Interest and Dividends Receivable Increase (Decrease) in Interest and Dividends Receivable, Total Cover [Abstract] Subsequent Event [Line Items] Net loss per share attributable to common stockholders, basic Earnings Per Share, Basic Earnings Per Share, Basic, Total Debt Securities, Available-for-Sale, Accumulated Gross Unrealized Gain, before Tax Unrealized Gains Operating expenses: Operating Expenses [Abstract] RSUs assumed for the Acquisition of AnHeart Options assumed for the Acquisition of AnHeart Stock Issued During Period, Shares, Acquisitions Contract liabilities, current portion Contract with Customer, Liability, Current Issuance of Replacement Awards Related to the AnHeart Acquisition, Total Issuance of replacement awards related to the AnHeart acquisition Issuance of Replacement Awards Related to the AnHeart Acquisition Issuance of Replacement Awards Related to the AnHeart Acquisition License revenue License [Member] Share-Based Payment Arrangement, Expense Stock-based compensation expense Amortization of premium on marketable securities. Amortization Of Premium On Marketable Securities Amortization of premium on marketable securities Schedule of Final Allocable Purchase Price Consists Schedule of Business Acquisitions, by Acquisition [Table Text Block] Accumulated Deficit Accumulated deficit Retained Earnings (Accumulated Deficit) Retained Earnings (Accumulated Deficit), Total Retained Earnings (Accumulated Deficit), Beginning Balance Retained Earnings (Accumulated Deficit), Ending Balance Accumulated deficit Plan Name Plan Name [Axis] Outstanding at June 30, 2024 Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Outstanding, Number Outstanding at December 31, 2023 Current assets: Assets, Current [Abstract] Common Stock, Capital Shares Reserved for Future Issuance Number of shares available for issuance Common stock reserved for future issuance Schedule Of Defined Benefit Plans Disclosures [Table] Defined Benefit Plan [Table] Net cash used in operating activities Net cash used in operating activities Net Cash Provided by (Used in) Operating Activities Net cash used in operating activities Product and Service [Domain] Accumulated other comprehensive (loss) income Accumulated Other Comprehensive Income (Loss), Net of Tax Accumulated Other Comprehensive Income (Loss), Net of Tax, Total Balance at beginning of period Balance at end of period Concentration Risk [Line Items] Number of common shares issuable under the plan (in shares) Number Of Shares Issuable Under Share Based Payment Arrangement Number of shares issuable under share based payment arrangement. Share-based compensation arrangement by share-based payment award, description Share-Based Compensation Arrangement by Share-Based Payment Award, Description Property and equipment, net of accumulated depreciation of \$782 and \$666, respectively Property, Plant and Equipment, Net Property, Plant and Equipment, Net, Total Property, Plant and Equipment, Net, Beginning Balance Property, Plant and Equipment, Net, Ending Balance 2025 2027 and thereafter Investment Securities Investment, Policy [Policy Text Block] Contract liabilities, net of current portion Contract with Customer, Liability, Noncurrent Cash flows from operating activities: Net Cash Provided by (Used in) Operating Activities [Abstract] Intangible assets, net of amortization of \$138 Intangible Assets Net of Amortization Intangible Assets Net of Amortization Common stock and additional paid in capital, par value (in Dollars per share) Common Stock, Par or Stated Value Per Share Sale of Stock, Description of Transaction Sale of Stock, Description of Transaction Contract Liability Contract with Customer, Liability Contract with Customer, Liability, Total Comprehensive loss: Comprehensive Income (Loss), Net of Tax, Attributable to Parent [Abstract] Non-cash lease expense. Non Cash Lease Expense Non-cash lease expense Statement of Cash Flows [Abstract] Antidilutive Securities, Name Antidilutive Securities, Name [Domain] Class of Warrant or Right, Number of Securities Called by Warrants or Rights Warrants to purchase common stock outstanding Selling, General and Administrative Expenses [Member] General and Administrative [Member] Long term debt

Long-Term Borrowing Long-Term Borrowing Equity Components Equity Components [Axis] Entity Address, Address Line One Entity Address, Address Line One Schedule of Concentration of Risk Concentration of customers Schedules of Concentration of Risk, by Risk Factor [Table Text Block] Computer [Member] Computer Equipment [Member] Equity Securities, FV-NI, Realized Gain Net realized gains on equities sold Borrowings Subordinated Borrowings Disclosure [Text Block] Subsequent Event Type [Axis] David Hanley [Member] David Hanley [Member] David Hanley [Member] Nonoperating Income (Expense) Total other income (expense), net Summary of Securities Outstanding Excluded from Calculation of Weighted Average Shares Outstanding Schedule of Antidilutive Securities Excluded from Computation of Earnings Per Share [Table Text Block] Share-based Compensation Arrangement by Share-based Payment Award, Fair Value Assumptions, Risk Free Interest Rate Risk-free interest rate U.S. Government And Government Agency Securities [Member] US Treasury and Government [Member] Subordinated Borrowings [Abstract] Amendment Flag Amendment Flag Operating Lease, Liability, Noncurrent Non-current operating lease liabilities Employee Stock Purchase Plan. Employee Stock Purchase Plan [Member] Corporate Bonds [Member] Corporate Debt Securities [Member] Short term working capital Short-Term Working Capital Short-Term Working Capital Sale of Stock, Price Per Share Stock price Strike price Net loss Options assumed for the acquisition of AnHeart Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Exercises in Period Exercised Exercise of stock options, shares Exercisable at June 30, 2024 Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Exercisable, Number Research and Development Expense [Member] Research and Development [Member] Equity Component Equity Component [Domain] Statistical Measurement Statistical Measurement [Domain] AnHeart [Member] AnHeart [Member] AnHeart [Member] Currency translation adjustment Currency translation adjustment Translation Adjustment Functional to Reporting Currency, Tax Expense (Benefit) Furniture and Fixtures [Member] Furniture and Equipment [Member] Innovent Innovent [Member] Innovent [Member] Net loss Net Income (Loss), Including Portion Attributable to Noncontrolling Interest Net Income (Loss), Including Portion Attributable to Noncontrolling Interest, Total Subordinated Borrowing [Table] Preferred Stock, Shares Issued Lease security deposit Operating Lease, Payments Proceeds from Stock Options Exercised Proceeds from exercises of options non-vested at June 30, 2024 Non-vested at December 31, 2023 Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Nonvested, Number of Shares Operating lease expense Operating Lease, Expense Prepaid expenses and other current assets Prepaid Expense and Other Assets, Current Preferred Class A [Member] Preferred Class A [Member] Collaborative Arrangement and Arrangement Other than Collaborative [Axis] Class of Stock Class of Stock [Domain] Other comprehensive loss Other Comprehensive Income (Loss), Net of Tax Other Comprehensive Income (Loss), Net of Tax, Total Comprehensive loss Comprehensive Income (Loss), Net of Tax, Attributable to Parent Common stock and additional paid in capital, shares outstanding Common Stock, Shares, Outstanding Common Stock, Shares, Outstanding, Beginning Balance Common Stock, Shares, Outstanding, Ending Balance Common stock outstanding Issuance of Common Stock related to the AnHeart acquisition Issuance of Common Stock Related to the AnHeart Acquisition Issuance of Common Stock Related to the AnHeart Acquisition Concentration of credit suppliers represent total purchases Concentration of Credit Suppliers Represent Total Purchases Concentration of Credit Suppliers Represent Total Purchases Preferred Stock, Par or Stated Value Per Share Preferred stock par value Warrant Disclosure [Abstract] Warrant disclosure. Revenue Benchmark [Member] Revenue Benchmark [Member] Private Placement Warrants [Member] Private Placement Warrants [Member] Private Placement Warrants member. Borrowing In 2022 [Member] Borrowing Two Thousands And Twenty Two [Member] Borrowing Two Thousands And Twenty Two [Member] Outstanding line of credit balance net Repayments of Lines of Credit Purchase [Member] Purchase [Member] Municipal Bonds [Member] Municipal Bonds [Member] Antidilutive Securities Antidilutive Securities [Axis] Debt Securities, Available-for-Sale, Realized Loss Realized losses from sales of available-for-sale securities Warrant Disclosure [Text Block] Warrant disclosure. Warrants Defined Benefit Plan Disclosure [Line Items] Defined Benefit Plan Disclosure [Line Items] Summary of Restricted Stock Units Activity Share-Based Payment Arrangement, Restricted Stock Unit, Activity [Table Text Block] Entity Interactive Data Current Entity Interactive Data Current Cash acquired from AnHeart acquisition Cash Acquired from Acquisition Line of Credit Facility, Frequency of Commitment Fee Payment Line of credit facility, frequency of commitment fee payment Business Acquisition, Equity Interest Issued or Issuable, Number of Shares Issuance of share pursuant to acquisition Document Quarterly Report Document Quarterly Report Related Party Transaction [Domain] Cash and cash equivalents, beginning of the period Cash and cash equivalents, end of the period Cash, Cash Equivalents, Restricted Cash, and Restricted Cash Equivalents Cash, Cash Equivalents, Restricted Cash and Restricted Cash Equivalents, Total Transaction costs related to AnHeart acquisition Transaction costs related to AnHeart acquisition Transaction costs related to AnHeart acquisition Statistical Measurement Statistical Measurement [Axis] Interest receivable on marketable securities Interest Receivable On Marketable Securities Current Interest receivable on marketable securities current. Cost of revenue Cost of Goods and Services Sold Cost of Goods and Services Sold, Total Operating Lease, Cost Operating Leases Number of Equity Securities Sold. Number of Equity Securities Sold Number of equity securities sold Schedule of revenue recognized Disaggregation of Revenue [Table Text Block] Schedule of Future Minimum Lease Commitments Schedule of Future Minimum Lease Commitments Amortization Amortization Amortization, Total Commercial Paper [Member] Commercial Paper [Member] Net cash consideration Payments to Acquire Businesses, Net of Cash Acquired Payments to Acquire Businesses, Net of Cash Acquired, Total Loss Contingency, Nature [Domain] Effect of foreign exchange rate changes on cash and cash equivalents Effect of Exchange Rate on Cash, Cash Equivalents, Restricted Cash, and Restricted Cash Equivalents, Continuing Operations Research and Development Costs Research and Development Expense, Policy [Policy Text Block] Increase (Decrease) in Prepaid Expense Prepaid expenses and other current assets Income Statement Location Statement of Income Location, Balance [Axis] Cash and Cash Equivalents [Axis] Total assets Assets Share-based Compensation Arrangement by Share-based Payment Award, Fair Value Assumptions, Expected Term Expected term in years Statement of Other Comprehensive Income [Abstract] Business Acquisition [Line Items] Business Acquisition [Line Items] Acquired in-process research and development Acquired In-Process Research and Development Acquired in-process research and development New Accounting Pronouncements, Policy [Policy Text Block] Accounting Pronouncements Not Yet Adopted Entity Address, City or Town Entity Address, City or Town Nonrecurring Adjustment [Domain] AOCI Attributable to Parent [Member] Accumulated Other Comprehensive Income [Member] Equity [Abstract] Long term debt maturity date Long-Term Debt, Maturity Date Share Price Due With in one year, Amortized Cost Debt Securities, Available-for-Sale, Amortized Cost, Maturity, Allocated and Single Maturity Date, Year One Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Outstanding, Period Increase (Decrease), Weighted Average Exercise Price Weighted average exercise price of outstanding options Share-Based Payment Arrangement [Text Block] Stock-Based Compensation Series A Preferred Stock Series A Preferred Stock [Member] Operating Lease, Weighted Average Discount Rate, Percent Weighted Average Discount Rate Percent Fair Value Disclosures [Abstract] Stock Issued During Period, Value, Stock Options Exercised Exercise of stock options Lessee, Lease, Description [Line Items] Current liabilities: Liabilities, Current [Abstract] Supplier B [Member] Supplier B [Member] Supplier B [Member] Fair Value Disclosures And Available For Sale Securities [Abstract] Fair Value Disclosures And Available For Sale Securities Abstract Public Warrants [Member] Public Warrants [Member] Public Warrants member Accounts Receivable [Member] Customer B [Member] Accounts Receivable [Member] Non-cash financing activity: Non-Cash Financing Activity [Abstract] Non-Cash Financing Activity [Abstract] Document Type Document Type Title of 12(b) Security Title of 12(b) Security US Government Debt Securities [Member] U.S. Government Securities [Member] Cash and cash equivalents Cash and Cash Equivalents, at Carrying Value Cash and Cash Equivalents, at Carrying Value, Total Cash and Cash Equivalents, at Carrying Value, Beginning Balance Cash and Cash Equivalents, at Carrying Value, Ending Balance 2027 and thereafter 2026 and thereafter Statement of Stockholders' Equity [Abstract] Borrowing In 2020 [Member] Borrowing Two Thousands And Twenty [Member] Borrowing Two Thousands And Twenty [Member] Document Fiscal Period Focus Document Fiscal Period Focus Customer B [Member] Customer B [Member] Customer B [Member] Research and development Research and Development Expense Research and Development Expense, Total Realized loss on marketable securities Realized loss on marketable securities Total losses on marketable securities Marketable Security, Gain (Loss) Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Grants in Period, Gross Granted CollaborationAndLicenseAgreementsTextBlock Collaboration And License Agreements TextBlock Collaboration And License Agreements TextBlock Principles of Consolidation Consolidation, Policy [Policy Text Block] Subsequent Event Type [Domain] Change in fair value of private warrants liability recognized in earnings. Change in Fair Value of Private Warrants Liability Recognized in Earnings Change in fair value of Private Warrants liability recognized in earnings Merger Agreement [Member] Merger agreement member. Lease Additional Term Lease additional term. Lease Additional Term Debt Securities, Available-for-Sale, Continuous Unrealized Loss Position, 12 Months or Longer 12 Months or Greater, Fair Value Class of warrant or right, exercise price of warrants or rights Class of Warrant or Right, Exercise Price of Warrants or Rights Financial Instruments Financial Instruments [Domain] Revenue, remaining performance obligation period Revenue, Remaining Performance Obligation, Expected Timing of Satisfaction, Period Class Of Warrant Or Right [Line Items] Class of Warrant or Right [Line Items] Cash flow from investing activities: Net Cash Provided by (Used in) Investing Activities [Abstract] Available-for-Sale Securities, Debt Maturities [Abstract] Liquidity Liquidity Policy [Text Block] Liquidity Policy Text Block. Marketable Securities, Current Marketable Securities, Current, Total Marketable securities Revenue Revenue from Contract with Customer, Excluding Assessed Tax Two thousand and twenty one Equity Incentive Plan and two thousand and nineteen equity incentive plan [Member] Two Thousand And Twenty One Equity Incentive Plan and Two Thousand And Nineteen Equity Incentive Plan [Member] 2021 and 2019 Equity Incentive Plan [Member] Lease security deposit Lease Security Deposit Noncurrent Lease security deposit noncurrent. Warrants and Rights Note Disclosure [Abstract] Selling, general and administrative Selling, General and Administrative Expense Selling, General and Administrative Expense, Total Business Combination, Asset Acquisition, and Joint Venture Formation [Abstract] Cash and cash equivalents and marketable securities Cash And Cash Equivalents And Marketable Securities Cash and cash equivalents and marketable securities . Intangible assets, net of amortization of \$138 Finite-Lived Intangible Assets, Net Finite-Lived Intangible Assets, Net, Ending Balance Finite-Lived Intangible Assets, Net, Beginning Balance Finite-Lived Intangible Assets, Net, Total Depreciation and amortization Depreciation, Depletion and Amortization, Nonproduction Depreciation, Depletion and Amortization, Nonproduction (Decrease) in Accounts Payable Increase (Decrease) in Accounts Payable (Decrease) in Accounts Payable Increase (Decrease) in Accounts Payable, Total Entity Address, Address Line Two Liabilities, mezzanine equity and stockholders' equity Liabilities and Equity [Abstract] Document Period End Date Document Period End Date Net cash provided by financing activities Net Cash Provided by (Used in) Financing Activities Common Class A [Member] Class A Common Stock [Member] Class A Shares [Member] Antidilutive Securities Excluded from Computation of Earnings Per Share, Amount Securities outstanding excluded from calculation of weighted average shares outstanding Stock-based compensation APIC, Share-Based Payment Arrangement, Increase for Cost Recognition APIC, Share-based Payment Arrangement, Increase for Cost Recognition, Total Common Class B [Member] Class B Common Stock [Member] Class B Shares [Member] Accrued expenses Accrued Liabilities, Current Accrued Liabilities, Current, Total Provision for income taxes Income Tax Expense (Benefit) Income Tax Expense (Benefit), Total Exchange Traded Funds [Member] Exchange Traded Funds [Member] Equity Interest Issued or Issuable, Type [Domain] Net loss on disposal of property and equipment Net loss on disposal of property and equipment Gain (Loss) on Disposition of Property Plant Equipment Gain (Loss) on Disposition of Property Plant Equipment, Total Trading Symbol Trading Symbol Business Combinations [Abstract] Equity, Attributable to Parent Total stockholders' equity Balance Balance Unaudited Interim Financial Information Unaudited Interim Financial Information Policy Policy [Text Block] Unaudited Interim Financial Information policy. Fixed interest rate on borrowings Subordinated Borrowing, Interest Rate Issuance Of Convertible Preferred Stocks Related To The Acquisition Of AnHeart(Shares) Issuance Of Convertible Preferred Stocks Related To The Acquisition Of AnHeart(Shares) Issuance Of Convertible Preferred Stocks Related To The Acquisition Of AnHeart(Shares) Issuance Of Replacement Awards Related To The AnHeart Acquisition Issuance Of Replacement Awards Related To The AnHeart Acquisition Value Issuance Of Replacement Awards Related To The AnHeart Acquisition Value Long-Lived Tangible Asset Long-Lived Tangible Asset [Axis] Class of Warrant or Right, Outstanding Aggregate warrants outstanding Warrant exercisable 2021 Equity Incentive Plan [Member] 2021 Equity Incentive Plan. Two Thousand And Twenty One Equity Incentive Plan [Member] Stock-based Compensation Share-Based Payment Arrangement [Policy Text Block] Restricted Stock Unit [Member] Restricted Stock Units (RSUs) [Member] Variable Lease, Cost Variable Lease Expenses Equity Securities, FV-NI, Gain (Loss), Total Equity Securities, FV-NI, Gain (Loss) Acquisition-related Costs [Member] Acquisition-related Costs [Member] Share-Based Payment Arrangement, Expensed and Capitalized, Amount [Table Text Block] Summary of Stock-Based Compensation Expense Nature of Operations Nature of Operations [Text Block] Geographical [Axis] Debt Securities, Available-for-Sale, Accumulated Gross Unrealized Loss, before Tax Unrealized Losses Leases [Abstract] Issuance of Common Stock for RSUs Vested, Share Issuance of Common Stock for RSUs Vested, Share Use of Estimates Use of Estimates, Policy [Policy Text Block] Cash and Cash Equivalents Cash and Cash Equivalents, Policy [Policy Text Block] Schedule Of Business Acquisitions By Acquisition [Table] Schedule of Business Acquisitions, by Acquisition [Table] Share-based Compensation Arrangement by Share-based Payment Award, Fair Value Assumptions, Expected Dividend Rate Dividend Schedule Of Contract Assets And Contract Liabilities Contract with Customer, Contract Asset, Contract Liability, and Receivable [Table Text Block] Security Exchange Name Security Exchange Name Schedule of Available-for-Sale Securities Reconciliation [Table Text Block] Schedule of Cost, Aggregate Fair Value and Unrealized Gains of Marketable Securities Available-for-sale Business Acquisition, Acquiree Business Acquisition, Acquiree [Domain] Redeemable Warrants [Member] Redeemable Warrants [Member] Total operating expenses Operating Expenses Shares based weighted average grant-date fair value of stock options Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Vested, Weighted Average Grant Date Fair Value Long-Lived Tangible Asset Long-Lived Tangible Asset [Domain] Loss Contingency Nature [Axis] Class of Stock Class of Stock [Axis] Property and equipment, net of accumulated depreciation of \$782 and \$666, respectively Accumulated Depreciation, Depletion and Amortization, Property, Plant, and Equipment Accumulated Depreciation, Depletion and Amortization, Property, Plant, and Equipment, Ending Balance Accumulated Depreciation, Depletion and Amortization, Property, Plant, and Equipment, Beginning Balance Accumulated Depreciation, Depletion and Amortization, Property, Plant, and Equipment, Total Fair Value Measurements and Marketable Securities Available for Sale Fair Value Disclosures [Text Block] 2024 2026 Preferred Nonconvertible Stock [Member] Subsequent Event [Member] Maximum [Member] Maximum [Member] Total liabilities, mezzanine equity and stockholders' equity Liabilities and Equity Operating Lease, Right-of-Use Asset Operating lease right-of-use assets Vested Vested Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Vested, Number of Shares Net loss per share attributable to common stockholders, diluted Earnings Per Share, Diluted Earnings Per Share, Diluted, Total Liabilities assumed Business Combination, Recognized Identifiable Assets Acquired and Liabilities Assumed, Liabilities Business Combination, Recognized Identifiable Assets Acquired and Liabilities Assumed, Liabilities, Total Entity Ex Transition Period Entity Ex Transition Period Total liabilities Liabilities Accounts receivable, net of allowance for credit loss of \$nil Accounts receivable Accounts Receivable, after Allowance for Credit Loss Accounts Receivable, after Allowance for Credit Loss, Total Fair Value Hierarchy and NAV Fair Value Hierarchy and NAV [Domain] Total current assets Assets, Current Debt Securities, Available-for-Sale, Realized Gain Realized gains from sales of available-for-sale securities Net cash provided by (used in) investing activities Net Cash Provided by (Used in) Investing Activities Share issued Stock Issued During Period, Shares, New Issues Issuance of common stock, shares Lessee, Operating Lease, Existence of Option to Extend [true false] Lease Existence Of Option To Extend Entity Filer Category Entity Filer Category Assumed Share Based Awards [Member] Assumed Share Based Awards [Member] Assumed Share Based Awards Member Currency Translation Adjustment Currency Translation Adjustment Value Currency Translation Adjustment Value Fair value of warrants, trading price Fair Value of Warrants, Trading Price Fair Value of Warrants, Trading Price Purchases of marketable securities Payments to Acquire Marketable Securities Payments to Acquire Marketable Securities, Total Other income (expense): Nonoperating Income (Expense) [Abstract] Current Fiscal Year End Date Current Fiscal Year End Date Property and Equipment Property, Plant and Equipment, Policy [Policy Text Block] Less: imputed interest Less: imputed interest Repayments of short term borrowing Repayments of Short-Term Debt Repayments of Short-Term Debt, Total Stock-based compensation Share-Based Payment Arrangement, Noncash Expense Share-based Payment Arrangement, Noncash Expense, Total Significant Risks and Uncertainties Unusual Risks and Uncertainties [Table Text Block] Related Party Transaction [Axis] Issuance of common stock for purchase under ESPP Issuance of common stock for purchase under ESPP, value Issuance of common stock for purchase under ESPP, value Entity Registrant Name Entity Registrant Name Debt Securities, Available-for-sale, Unrealized Loss Position, Accumulated Loss, Total Debt Securities, Available-for-Sale, Unrealized Loss Position, Accumulated Loss Debt Securities, Available-for-sale, Unrealized Loss Position, Accumulated Loss Debt Securities, Available-for-sale, Unrealized Loss Position, Accumulated Loss Debt Securities, Available-for-Sale, Amortized Cost [Abstract] Estimated Fair Value Warrant Estimated Fair Value Warrant Per Share Estimated Fair Value Warrant Per Share Short term borrowings drawn down Short-Term Borrowings Drawn Down Short-Term Borrowings Drawn Down Gross Profit Gross profit Gross Profit Schedule Of Share Based Compensation Arrangements By Share Based Payment Award [Table] Schedule of Share-Based Compensation Arrangements by Share-Based Payment Award [Table] Net Loss per Share Earnings Per Share [Text Block] Entity Emerging Growth Company Entity Emerging Growth Company Revenue, performance obligation satisfied over time, method used, description Revenue, Performance Obligation Satisfied over Time, Method Used, Description Money Market Funds [Member] Money Market Funds [Member] Weighted average common shares outstanding, basic Weighted Average Number of Shares

Outstanding, Basic Weighted Average Number of Shares Outstanding, Basic, Total Marketable Securities [Abstract] Marketable securities: Standby Letters of Credit [Member] Standby Letter of Credit [Member] Loss Contingencies [Line Items] Loss Contingencies [Line Items] Business Acquisition, Date of Acquisition Agreement Agreement Date NK NK [Member] NK [Member] Common stock and additional paid in capital, shares authorized Common Stock, Shares Authorized Property, Plant and Equipment, Useful Life Property and equipment, estimated useful life Proceeds from sale of marketable securities Proceeds from Sale and Maturity of Marketable Securities Proceeds from Sale and Maturity of Marketable Securities, Total Concentration Risk Type [Axis] Cash and Cash Equivalents [Abstract] Cash equivalents: Level 1 [Member] Fair Value, Inputs, Level 1 [Member] Financial Instrument Financial Instrument [Axis] Other Comprehensive Income, Noncontrolling Interest [Text Block] Accumulated Other Comprehensive Income Share-Based Payment Arrangement [Abstract] Impairment of Long-Lived Assets Impairment or Disposal of Long-Lived Assets, Policy [Policy Text Block] Collaboration and Licence Agreement [Abstract] Collaboration and Licence Agreement [Abstract] Subordinated Borrowing, Name [Domain] Forward Purchase Warrants [Member] Forward Purchase Warrants [Member] Forward Purchase Warrants member. Issuance Of Common Stock For AnHeart Acquisition Issuance Of Common Stock For AnHeart Acquisition Value Issuance Of Common Stock For AnHeart Acquisition Value Schedule of Accumulated Other Comprehensive Income (Loss) [Table Text Block] Schedule of Changes in Accumulated Other Comprehensive Income Schedule Of Available for sale Securities Maturity On Fair Value Schedule Of Available For Sale Securities Classified By Contractual Maturity Date Table [Text Block] Schedule Of Available for sale Securities Classified By Contractual Maturity Date Table Text Block Debt Securities, Available-for-Sale Debt Securities, Available-for-sale, Total Fair Value Accounting Policies [Abstract] Amount reclassified for realized gain on marketable securities Realized loss On Marketable Securities Realized loss On Marketable Securities Amount reclassified for realized loss (gain) on marketable securities Share-Based Payment Arrangement, Option, Activity [Table Text Block] Summary of Stock Option Activity under 2019 Plan and 2021 Plan Long-Term Line of Credit Long-term Line of Credit, Total Letter of Credit, Amount Equity, Attributable to Parent [Abstract] Stockholders' equity Redeemable convertible preferred stocks [Member] Redeemable Convertible Preferred Stocks [Member] Redeemable Convertible Preferred Stocks [Member] Consideration Warrants [Member] Consideration Warrants [Member] Consideration Warrants [Member] Proceeds from issuance of Common Stock under ESPP Proceeds from Issuance of Common Stock under ESPP Proceeds from Issuance of Common Stock Under ESPP Short-term borrowings outstanding balance Short-Term Debt, Average Outstanding Amount Loan Agreements With China Merchants Bank [Member] Loan Agreements With China Merchants Bank [Member] Loan Agreements With China Merchants Bank [Member] Interest income Interest Income, Other Statement [Table] Statement [Table] Segment Information Segment Reporting, Policy [Policy Text Block] Subsequent Event [Table] Warrant liability. Warrant Liability Warrant liability Equity Interest Type [Axis] Fair Value Adjustment of Warrants Change in fair value of warrant liability Fair value of Warrants Change in fair value of warrant liability Revenue from customer Revenue From Contract With Customer Including Assessed Tax Revenue from Contract with Customer, Including Assessed Tax Entity File Number Entity File Number Issuance Of Common Stock For RSUs Vested (Shares) Issuance Of Common Stock For RSUs Vested (Shares) Issuance of Common Stock for RSUs Vested (Shares) Net decrease in cash and cash equivalents Cash, Cash Equivalents, Restricted Cash, and Restricted Cash Equivalents, Period Increase (Decrease), Including Exchange Rate Effect Net loss per share attributable to common stockholders, basic and diluted Earnings Per Share [Abstract] Subordinated Borrowing [Axis] Cash Equivalents [Member] Cash Equivalents [Member] Statement of Financial Position [Abstract] Interest expense Interest expense Interest Expense, Operating and Nonoperating Interest Expense, Operating and Nonoperating, Total Loss from operations Operating Income (Loss) Loss from operations Research and development service revenue Research and Development Service Revenue [Member] Research and Development Service Revenue [Member] Shares issued, price per share Shares Issued, Price Per Share Share-based Compensation Arrangement by Share-based Payment Award, Fair Value Assumptions, Expected Volatility Rate Expected volatility Common Stock, No Par Value Stock par value Changes in liabilities classified in Level 3 of fair value hierarchy. Changes in Liabilities Classified in Level 3 of Fair Value Hierarchy Beginning balance Ending balance Entity Shell Company Entity Shell Company Fair Value Measurement Inputs and Valuation Techniques [Table Text Block] Schedule of Quantitative Information Regarding Level 3 Fair Value Measurements Entity Tax Identification Number Entity Tax Identification Number Assets Assets [Abstract] Class of Warrant or Right Class of Warrant or Right [Domain] CHINA China [Member] Entity Current Reporting Status Entity Current Reporting Status Fair Value, Liabilities Measured on Recurring Basis [Table Text Block] Summary of Changes in liabilities classified in Level 3 of the fair value hierarchy Equity Securities, FV-NI, Realized Gain (Loss) [Abstract] Net recognized gains (losses) on equity securities were as follows: Subsequent Events [Abstract] Costs to Fulfill a Contract with a Customer Revenue from Contract with Customer [Policy Text Block] Customer A [Member] Customer A [Member] Customer A [Member] Assets acquired Business Combination, Recognized Identifiable Assets Acquired and Liabilities Assumed, Assets Business Combination, Recognized Identifiable Assets Acquired and Liabilities Assumed, Assets, Total Geographical [Domain] Document Fiscal Year Focus Document Fiscal Year Focus Certificate of deposits [Member] Certificate of deposits [Member] Certificate of deposits [Member] Class A convertible preferred stock, \$0.001 par value per share; 851,202 shares issued and outstanding as of June 30, 2024. Preferred Stock, Value, Issued Total, Amortized Cost Debt Securities, Available-for-Sale, Maturity, Allocated and Single Maturity Date, Amortized Cost Due after one year through five years, Amortized Cost Debt Securities, Available-for-Sale, Amortized Cost, Maturity, Allocated and Single Maturity Date, after Year One Through Five Business Acquisition Business Acquisition [Axis] City Area Code City Area Code Total current liabilities Liabilities, Current Organization, Consolidation and Presentation of Financial Statements [Abstract] Local Phone Number Local Phone Number Preferred Stock Shares Outstanding Preferred Stock, Shares Outstanding Preferred Stock, Shares Outstanding, Ending Balance Preferred Stock, Shares Outstanding, Beginning Balance Antidilutive Securities Excluded From Computation Of Earnings Per Share [Line Items] Antidilutive Securities Excluded from Computation of Earnings Per Share [Line Items] Debt Securities, Available-for-Sale, Continuous Unrealized Loss Position, Less than 12 Months, Accumulated Loss Less than 12 Months, Unrealized Losses Issuance of common stock for purchase under ESPP, (shares) Issuance of common stock for purchase under ESPP, (shares) Issuance of common stock for purchase under ESPP, shares Income Tax, Policy [Policy Text Block] Income Taxes Commitments and Contingencies Disclosure [Abstract] Series A Non-Voting Convertible Preferred Stock [Member] Convertible Preferred Stock [Member] Convertible Preferred Stock [Member] Financial Liabilities Fair Value Disclosure [Abstract] Financial liabilities: Debt Securities, Available-for-Sale, Amortized Cost Amortized Cost Investment advisory fees Professional Fees Operating Lease, Liability, Current Current operating lease liabilities Document Information [Line Items] Jerry Wang, AnHeart's [Member] Jerry Wang, AnHeart's [Member] Jerry Wang, AnHeart's [Member] Jerry Wang, AnHeart's [Member] Shares issued upon conversion Convertible Preferred Stock, Shares Issued upon Conversion Retained Earnings [Member] Accumulated Deficit [Member] Document Transition Report Document Transition Report Kerry Wentworth [Member] Kerry Wentworth [Member] Kerry Wentworth [Member] Share-Based Compensation Arrangement by Share-Based Payment Award, Purchase Price of Common Stock, Percent Number of common stock outstanding or issuable, percentage 2023 2025 Schedule Of Antidilutive Securities Excluded From Computation Of Earnings Per Share [Table] Antidilutive Security, Excluded EPS Calculation [Table] Loan Agreements With Bank of Hangzhou [Member] Loan Agreements With Bank of Hangzhou [Member] Loan Agreements With Bank of Hangzhou [Member] Concentration of Risk Concentration Risk, Credit Risk, Policy [Policy Text Block] Minimum [Member] Minimum [Member] Minimum [Member] Common Stock [Member] Employee Service Share Based Compensation Allocation Of Recognized Period Costs [Line Items] Share-Based Payment Arrangement, Expensed and Capitalized, Amount [Line Items] Increase (Decrease) in Other Noncurrent Assets Other non-current assets Other non-current assets Issuance of warrants related to the AnHeart acquisition Issuance of Warrants Related to the AnHeart Acquisition Issuance of Warrants Related to the AnHeart Acquisition Additional Paid-in Capital [Member] Additional Paid-in Capital [Member] Proceeds from borrowings Proceeds from Short-Term Debt Proceeds from Short-Term Debt, Total Functional Currency and Foreign Currency Translations Foreign Currency Transactions and Translations Policy [Policy Text Block] Adjustments to reconcile net loss to net cash used in operating activities: Adjustments to Reconcile Net Income (Loss) to Cash Provided by (Used in) Operating Activities [Abstract] Other comprehensive loss, net of taxes: Other Comprehensive Income (Loss), Net of Tax [Abstract] Income Statement Location Statement of Income Location, Balance [Domain] 2024 (remaining 6 months) 2022 (remaining 9 months) Other Assets, Noncurrent Other non-current assets Revenue Recognition Revenue Recognition Policy [Policy Text Block] Revenue Recognition Policy [Policy Text Block] Issuance of Convertible Preferred Stocks Related to The Acquisition of AnHeart Issuance of Convertible Preferred Stocks Related to The Acquisition of AnHeart Value Issuance of Convertible Preferred Stocks Related to The Acquisition of AnHeart Value Concentration Risk, Percentage Concentration Risk Threshold Percentage Concentration Risk Threshold Percentage Payments on borrowings Payments on borrowings Payments of Financing Costs Payments of Financing Costs, Total Amount reclassified for realized loss (gain) on marketable securities Amount reclassified for realized loss (gain) on marketable securities Debt Securities, Available-for-sale, Unrealized Gain (Loss), Total Debt Securities, Available-for-Sale, Unrealized Gain (Loss) Entity Common Stock, Shares Outstanding Entity Common Stock, Shares Outstanding Forfeited or canceled Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Nonvested Options Forfeited, Number of Shares Change in unrealized (loss) gain on available-for-sale securities Debt Securities, Unrealized Gain (Loss), Total Debt Securities, Unrealized Gain (Loss) Debt Securities, Available-for-Sale, Continuous Unrealized Loss Position, Less than 12 Months Less than 12 Months, Fair Value Basis of Presentation Basis of Accounting, Policy [Policy Text Block] Total allocable purchase price Allocable Purchase Price Allocable Purchase Price Warrants and Rights Outstanding Fair values of outstanding warrants Class A and Class B common stock and additional paid in capital, \$0.0001 par value per share; 1,060,000,000 (Class A 1,000,000,000, Class B 60,000,000) shares authorized as of June 30, 2024 and December 31, 2023, 248,624,729 (Class A 247,624,729, Class B 1,000,000) and 219,046,219 (Class A 218,046,219, Class B 1,000,000) shares issued and outstanding as of June 30, 2024 and December 31, 2023, respectively Common Stock, Value, Issued Common Stock, Value, Issued, Total Common Stock, Value, Issued, Beginning Balance Common Stock, Value, Issued, Ending Balance Convertible Preferred Stocks [Member] Convertible Preferred Stocks [Member] Convertible Preferred Stocks [Member] Fair Value, Assets Measured on Recurring Basis [Table Text Block] Schedule of Marketable securities Measured at Fair Value on Recurring Basis Issuance of Common Stock for AnHeart acquisition (Shares) Issuance of Common Stock for AnHeart acquisition (Shares) Issuance of Common Stock for AnHeart acquisition (Shares) Total operating lease liabilities Total operating lease liabilities Total operating lease liabilities Commitments and contingencies (Note 12) Commitments and Contingencies Warrants Policy [Text Block] Warrants policy text block. Warrants Debt Securities, Available-for-Sale, Continuous Unrealized Loss Position, 12 Months or Longer, Accumulated Loss 12 Months or Greater, Unrealized Losses Weighted average common shares outstanding, diluted Weighted Average Number of Shares Outstanding, Diluted Weighted Average Number of Shares Outstanding, Diluted Total Significant Accounting Policies Significant Accounting Policies [Text Block] Contract liabilities Increase (Decrease) in Commodity Contract Assets and Liabilities Lessee, Operating Lease, Liability, to be Paid Total undiscounted lease payments Leases policy text block. Leases [Policy Text Block] Leases Concentration Risk Benchmark [Domain] Gary Hattersley [Member] Gary Hattersley [Member] Gary Hattersley [Member] Entity Small Business Entity Small Business Entity Bankruptcy Proceedings, Reporting Current Entity Bankruptcy Proceedings, Reporting Current Revenues Revenues, Total Cash flow from financing activities: Net Cash Provided by (Used in) Financing Activities [Abstract] Mezzanine equity; Mezzanine Equity [Abstract] Mezzanine Equity [Abstract] Short term bank loan Broker-Dealer, Bank Loan, Short-Term Level 3 [Member] Fair Value, Inputs, Level 3 [Member] Operating Lease, Weighted Average Remaining Lease Term Weighted Average Remaining Lease Term Subsequent Events [Text Block] Subsequent Event Accounts payable Accounts Payable, Current Accounts Payable, Current, Total Total losses on marketable securities Realized loss on marketable securities Marketable Securities, Gain (Loss) Risks and Uncertainties [Abstract] Class B Common Shares Subject to Repurchase. Class B Common Shares Subject to Repurchase [Member] Accrued expenses Increase (Decrease) in Accrued Liabilities Increase (Decrease) in Accrued Liabilities, Total Accounts receivable, net of allowance for credit loss Accounts Receivable, Allowance for Credit Loss Accounts Receivable, Allowance for Credit Loss, Ending Balance Accounts Receivable, Allowance for Credit Loss, Beginning Balance XML 8 R1.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Document and Entity Information - shares 6 Months Ended Jun. 30, 2024 Jul. 29, 2024 Document Information [Line Items] A A Document Type 10-Q A Amendment Flag false A Entity Registrant Name Nuvation Bio Inc. A Document Fiscal Year Focus 2024 A Document Period End Date Jun. 30, 2024 A Document Fiscal Period Focus Q2 A Current Fiscal Year End Date -12-31 A Entity Central Index Key 0001811063 A Document Quarterly Report true A Document Transition Report false A Entity File Number 001-39351 A Entity Incorporation, State or Country Code DE A Entity Tax Identification Number 85-0862255 A Entity Address, Address Line One 1500 Broadway A Entity Address, Address Line Two Suite 1401 A Entity Address, City or Town New York A Entity Address, State or Province NY A Entity Address, Postal Zip Code 10036 A Local Phone Number 208-6102 A City Area Code 332 A Entity Current Reporting Status Yes A Entity Interactive Data Current Yes A Entity Filer Category Non-accelerated Filer A Entity Small Business true A Entity Emerging Growth Company false A Entity Shell Company false A Entity Common Stock, Shares Outstanding A 249,245,129 Class A Common Stock [Member] A A Document Information [Line Items] A A Title of 12(b) Security Class A Common Stock, \$0.0001 par value per share A Trading Symbol NUVB A Security Exchange Name NYSE A Redeemable Warrants [Member] A A Document Information [Line Items] A A Title of 12(b) Security Redeemable Warrants, each whole warrantexercisable for one share of Common Stock at anexercise price of \$11.50 per share A Trading Symbol NUVB.WS A Security Exchange Name NYSE A X - DefinitionBoolean flag that is true when the XBRL content amends previously-filed or accepted submission. + ReferencesNo definition available. + Details Name: dei_AmendmentFlag Namespace Prefix: dei_Data Type: xbrli:booleanItemType Balance Type: na Period Type: duration X - DefinitionArea code of city + ReferencesNo definition available. + Details Name: dei_CityAreaCode Namespace Prefix: dei_Data Type: xbrli:normalizedStringItemType Balance Type: na Period Type: duration X - DefinitionEnd date of current fiscal year in the format --MM-DD. + ReferencesNo definition available. + Details Name: dei_CurrentFiscalYearEndDate Namespace Prefix: dei_Data Type: xbrli:gMonthDayItemType Balance Type: na Period Type: duration X - DefinitionFiscal period values are FY, Q1, Q2, and Q3. 1st, 2nd and 3rd quarter 10-Q or 10-QT statements have value Q1, Q2, and Q3 respectively, with 10-K, 10-KT or other fiscal year statements having FY. + ReferencesNo definition available. + Details Name: dei_DocumentFiscalPeriodFocus Namespace Prefix: dei_Data Type: dei:fiscalPeriodItemType Balance Type: na Period Type: duration X - DefinitionThis is focus fiscal year of the document report in YYYY format. For a 2006 annual report, which may also provide financial information from prior periods, fiscal 2006 should be given as the fiscal year focus. Example: 2006. + ReferencesNo definition available. + Details Name: dei_DocumentFiscalYearFocus Namespace Prefix: dei_Data Type: xbrli:gYearItemType Balance Type: na Period Type: duration X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesNo definition available. + Details Name: dei_DocumentInformationLineItems Namespace Prefix: dei_Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionFor the EDGAR submission types of Form 8-K: the date of the report, the date of the earliest event reported; for the EDGAR submission types of Form N-1A: the filing date; for all other submission types: the end of the reporting or transition period. The format of the date is YYYY-MM-DD. + ReferencesNo definition available. + Details Name: dei_DocumentPeriodEndDate Namespace Prefix: dei_Data Type: xbrli:dateTimeItemType Balance Type: na Period Type: duration X - DefinitionBoolean flag that is true only for a form used as an quarterly report. + ReferencesReference 1: http://www.xbrl.org/2003/role/presentationRef -Publisher SEC -Name Form 10-Q -Number 240 -Section 308 -Subsection a + Details Name: dei_DocumentQuarterlyReport Namespace Prefix: dei_Data Type: xbrli:booleanItemType Balance Type: na Period Type: duration X - DefinitionBoolean flag that is true only for a form used as a transition report. + ReferencesReference 1: http://www.xbrl.org/2003/role/presentationRef -Publisher SEC -Name Forms 10-K, 10-Q, 20-F -Number 240 -Section 13 -Subsection a-1 + Details Name: dei_DocumentTransitionReport Namespace Prefix: dei_Data Type: xbrli:booleanItemType Balance Type: na Period Type: duration X - DefinitionThe type of document being provided (such as 10-K, 10-Q, 485BPOS, etc). The document type is limited to the same value as the supporting SEC submission type, or the word 'Other'. + ReferencesNo definition available. + Details Name: dei_DocumentType Namespace Prefix: dei_Data Type: dei:submissionItemType Balance Type: na Period Type: duration X - DefinitionAddress Line 1 such as Attn, Building Name, Street Name + ReferencesNo definition available. + Details Name: dei_EntityAddressAddressLine1 Namespace Prefix: dei_Data Type: xbrli:normalizedStringItemType Balance Type: na Period Type: duration X - DefinitionAddress Line 2 such as Street or Suite number + ReferencesNo definition available. + Details Name: dei_EntityAddressAddressLine2 Namespace Prefix: dei_Data Type: xbrli:normalizedStringItemType Balance Type: na Period Type: duration X - DefinitionName of the City or Town + ReferencesNo definition available. + Details Name: dei_EntityAddressCityOrTown Namespace Prefix: dei_Data Type: xbrli:normalizedStringItemType Balance Type: na Period Type: duration X - DefinitionCode for the postal or zip code + ReferencesNo definition available. + Details Name: dei_EntityAddressPostalZipCode Namespace Prefix: dei_Data Type: xbrli:normalizedStringItemType Balance Type: na Period Type: duration X - DefinitionName of the state or province. + ReferencesNo definition available. + Details Name: dei_EntityAddressStateOrProvince Namespace Prefix: dei_Data Type: dei:stateOrProvinceItemType Balance Type: na Period Type: duration X - DefinitionA unique 10-digit SEC-issued value to identify entities that have filed disclosures with the SEC. It is commonly abbreviated as CIK. + ReferencesReference 1: http://www.xbrl.org/2003/role/presentationRef -Publisher SEC -Name Exchange Act -Number 240 -Section 12 -Subsection b-2 + Details Name:

dei_EntityCentralIndexKey Namespace Prefix: dei_ Data Type: dei:centralIndexKeyItemType Balance Type: na Period Type: duration X - DefinitionIndicate number of shares or other units outstanding of each of registrant's classes of capital or common stock or other ownership interests, if and as stated on cover of related periodic report. Where multiple classes or units exist define each class/interest by adding class of stock items such as Common Class A [Member], Common Class B [Member] or Partnership Interest [Member] onto the Instrument [Domain] of the Entity Listings, Instrument. + ReferencesNo definition available. + Details Name: dei_EntityCommonStockSharesOutstanding Namespace Prefix: dei_ Data Type: xbrli:sharesItemType Balance Type: na Period Type: instant X - DefinitionIndicate 'Yes' or 'No' whether registrants (1) have filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that registrants were required to file such reports), and (2) have been subject to such filing requirements for the past 90 days. This information should be based on the registrant's current or most recent filing containing the related disclosure. + ReferencesNo definition available. + Details Name: dei_EntityCurrentReportingStatus Namespace Prefix: dei_ Data Type: dei:yesNoItemType Balance Type: na Period Type: duration X - DefinitionIndicate if registrant meets the emerging growth company criteria. + ReferencesReference 1: <http://www.xbrl.org/2003/role/presentationRef> -Publisher SEC -Name Exchange Act -Number 240 -Section 12 -Subsection b-2 + Details Name: dei_EntityEmergingGrowthCompany Namespace Prefix: dei_ Data Type: xbrli:booleanItemType Balance Type: na Period Type: duration X - DefinitionCommission file number. The field allows up to 17 characters. The prefix may contain 1-3 digits, the sequence number may contain 1-8 digits, the optional suffix may contain 1-4 characters, and the fields are separated with a hyphen. + ReferencesNo definition available. + Details Name: dei_EntityFileNumber Namespace Prefix: dei_ Data Type: dei:fileNumberItemType Balance Type: na Period Type: duration X - DefinitionIndicate whether the registrant is one of the following: Large Accelerated Filer, Accelerated Filer, Non-accelerated Filer. Definitions of these categories are stated in Rule 12b-2 of the Exchange Act. This information should be based on the registrant's current or most recent filing containing the related disclosure. + ReferencesReference 1: <http://www.xbrl.org/2003/role/presentationRef> -Publisher SEC -Name Exchange Act -Number 240 -Section 12 -Subsection b-2 + Details Name: dei_EntityFilerCategory Namespace Prefix: dei_ Data Type: dei:filerCategoryItemType Balance Type: na Period Type: duration X - DefinitionTwo-character EDGAR code representing the state or country of incorporation. + ReferencesNo definition available. + Details Name: dei_EntityIncorporationStateCountryCode Namespace Prefix: dei_ Data Type: dei:edgarStateCountryItemType Balance Type: na Period Type: duration X - DefinitionBoolean flag that is true when the registrant has submitted electronically every Interactive Data File required to be submitted pursuant to Rule 405 of Regulation S-T during the preceding 12 months (or for such shorter period that the registrant was required to submit such files). + ReferencesReference 1: <http://www.xbrl.org/2003/role/presentationRef> -Publisher SEC -Name Regulation S-T -Number 232 -Section 405 + Details Name: dei_EntityInteractiveDataCurrent Namespace Prefix: dei_ Data Type: dei:yesNoItemType Balance Type: na Period Type: duration X - DefinitionThe exact name of the entity filing the report as specified in its charter, which is required by forms filed with the SEC. + ReferencesReference 1: <http://www.xbrl.org/2003/role/presentationRef> -Publisher SEC -Name Exchange Act -Number 240 -Section 12 -Subsection b-2 + Details Name: dei_EntityRegistrantName Namespace Prefix: dei_ Data Type: xbrli:normlizedStringItemType Balance Type: na Period Type: duration X - DefinitionBoolean flag that is true when the registrant is a shell company as defined in Rule 12b-2 of the Exchange Act. + ReferencesReference 1: <http://www.xbrl.org/2003/role/presentationRef> -Publisher SEC -Name Exchange Act -Number 240 -Section 12 -Subsection b-2 + Details Name: dei_EntityShellCompany Namespace Prefix: dei_ Data Type: xbrli:booleanItemType Balance Type: na Period Type: duration X - DefinitionIndicates that the company is a Smaller Reporting Company (SRC). + ReferencesReference 1: <http://www.xbrl.org/2003/role/presentationRef> -Publisher SEC -Name Exchange Act -Number 240 -Section 12 -Subsection b-2 + Details Name: dei_EntitySmallBusiness Namespace Prefix: dei_ Data Type: xbrli:booleanItemType Balance Type: na Period Type: duration X - DefinitionThe Tax Identification Number (TIN), also known as an Employer Identification Number (EIN), is a unique 9-digit value assigned by the IRS. + ReferencesReference 1: <http://www.xbrl.org/2003/role/presentationRef> -Publisher SEC -Name Exchange Act -Number 240 -Section 12 -Subsection b-2 + Details Name: dei_EntityTaxIdentificationNumber Namespace Prefix: dei_ Data Type: dei:employerIdItemType Balance Type: na Period Type: duration X - DefinitionLocal phone number for entity. + ReferencesNo definition available. + Details Name: dei_LocalPhoneNumber Namespace Prefix: dei_ Data Type: xbrli:normlizedStringItemType Balance Type: na Period Type: duration X - DefinitionTitle of a 12(b) registered security. + ReferencesReference 1: <http://www.xbrl.org/2003/role/presentationRef> -Publisher SEC -Name Exchange Act -Number 240 -Section 12 -Subsection b + Details Name: dei_Security12bTitle Namespace Prefix: dei_ Data Type: dei:securityTitleItemType Balance Type: na Period Type: duration X - DefinitionName of the Exchange on which a security is registered. + ReferencesReference 1: <http://www.xbrl.org/2003/role/presentationRef> -Publisher SEC -Name Exchange Act -Number 240 -Section 12 -Subsection d1-1 + Details Name: dei_SecurityExchangeName Namespace Prefix: dei_ Data Type: dei:edgarExchangeCodeItemType Balance Type: na Period Type: duration X - DefinitionTrading symbol of an instrument as listed on an exchange. + ReferencesNo definition available. + Details Name: dei_TradingSymbol Namespace Prefix: dei_ Data Type: dei:tradingSymbolItemType Balance Type: na Period Type: duration X - Details Name: us-gaap_StatementClassOfStockAxis=us-gaap_CommonClassAMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_StatementClassOfStockAxis=nuvb_RedeemableWarrantsMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 9 R2.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Consolidated Balance Sheets (Unaudited) - USD (\$) \$ in Thousands Jun. 30, 2024 Dec. 31, 2023 Current assets: \$ 34,285 \$ 42,649 Accounts receivable, net of allowance for credit loss of \$11 17 0 Prepaid expenses and other current assets 5,991 1,519 Marketable securities 542,884 568,564 Interest receivable on marketable securities 3,895 3,702 Total current assets 587,172 616,434 Property and equipment, net of accumulated depreciation of \$782 and \$666, respectively 751 717 Intangible assets, net of amortization of \$138 2,932 0 Lease security deposit 143 141 Operating lease right-of-use assets 2,723 3,605 Other non-current assets 1,075 587 Total assets 594,796 621,484 Current liabilities: \$ 594,796 \$ 621,484 X - Accounts payable 5,211 2,209 Current operating lease liabilities 2,076 1,972 Contract liabilities, current portion 12,200 0 Short-term borrowings 11,634 0 Accrued expenses 19,974 9,793 Total current liabilities 51,095 13,974 Warrant liability 1,441 353 Contract liabilities, net of current portion 9,157 0 Non-current operating lease liabilities 972 2,035 Total liabilities 62,665 16,362 Commitments and contingencies (Note 12) Mezzanine equity: \$ 4 Class A convertible preferred stock, \$0.0001 par value per share; 851,202 shares issued and outstanding as of June 30, 2024, 274,938 0 Total Mezzanine Equity, Total 274,938 0 Stockholders' equity \$ 4 Class A and Class B common stock and additional paid in capital, \$0.0001 par value per share; 1,060,000,000 (Class A) 1,000,000,000, Class B 60,000,000 shares authorized as of June 30, 2024 and December 31, 2023, 248,624,729 (Class A) 247,624,729, Class B 1,000,000,000 and 219,046,219 (Class A) 218,046,219, Class B 1,000,000,000 shares issued and outstanding as of June 30, 2024 and December 31, 2023, respectively 1,078,547 947,745 Accumulated deficit (820,088) (342,804) Accumulated other comprehensive (loss) income (1,266) 181 Total stockholders' equity 257,193 605,122 Total liabilities, mezzanine equity and stockholders' equity \$ 594,796 \$ 621,484 X - DefinitionInterest receivable on marketable securities current. + ReferencesNo definition available. + Details Name: nuvb_InterestReceivableOnMarketableSecuritiesCurrent Namespace Prefix: nuvb_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionLease security deposit noncurrent. + ReferencesNo definition available. + Details Name: nuvb_LeaseSecurityDepositNoncurrent Namespace Prefix: nuvb_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionMezzanine Equity [Abstract] + ReferencesNo definition available. + Details Name: nuvb_MezzanineEquityAbstract Namespace Prefix: nuvb_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionTotal Mezzanine Equity + ReferencesNo definition available. + Details Name: nuvb_TotalMezzanineEquity Namespace Prefix: nuvb_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionWarrant liability. + ReferencesNo definition available. + Details Name: nuvb_WarrantLiability Namespace Prefix: nuvb_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionCarrying value as of the balance sheet date of liabilities incurred (and for which invoices have typically been received) and payable to vendors for goods and services received that are used in an entity's business. Used to reflect the current portion of the liabilities (due within one year or within the normal operating cycle if longer). + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(19)(a)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1>Reference 2: <http://www.xbrl.org/2003/role/exampleRef> -Topic 852 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 10 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481372/852-10-55-10> + Details Name: us-gaap_AccountsPayableCurrent Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionAmount, after allowance for credit loss, of right to consideration from customer for product sold and service rendered in normal course of business. + ReferencesReference 1: <http://www.xbrl.org/2003/role/exampleRef> -Topic 946 -SubTopic 31 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Subparagraph (d) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147477802/946-310-45-1>Reference 2: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(5)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478777/944-210-S99-1>Reference 3: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.6-04(5)(b)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479170/946-210-S99-1>Reference 4: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 954 -SubTopic 31 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479196/954-310-45-1>Reference 5: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(3)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478777/944-210-S99-1> + Details Name: us-gaap_AccountsReceivableNet Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionCarrying value as of the balance sheet date of obligations incurred and payable, pertaining to costs that are statutory in nature, are incurred on contractual obligations, or accumulate over time and for which invoices have not yet been received or will not be rendered. Examples include taxes, interest, rent and utilities. Used to reflect the current portion of the liabilities (due within one year or within the normal operating cycle if longer). + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(20)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1> + Details Name: us-gaap_AccruedLiabilitiesCurrent Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionAmount, after tax, of accumulated increase (decrease) in equity from transaction and other event and circumstance from nonowner source. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Topic 220 -SubTopic 10 -Section 45 -Paragraph 14A -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482790/220-10-45-14A>Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 11 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482790/220-10-45-11>Reference 3: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 944 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 2 -Subparagraph (g)(2)(ii) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480016/944-40-65-2>Reference 4: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 944 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 2 -Subparagraph (h)(2) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480016/944-40-65-2>Reference 5: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(30)(a)(4)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1>Reference 6: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(23)(a)(3)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478777/944-210-S99-1>Reference 7: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Topic 220 -SubTopic 10 -Section 45 -Paragraph 14 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482790/220-10-45-14> + Details Name: us-gaap_AccumulatedOtherComprehensiveIncomeLossNetOfTax Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionAmount of asset recognized for present right to economic benefit. + ReferencesReference 1: <http://www.xbrl.org/2003/role/exampleRef> -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 48 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482785/280-10-55-48>Reference 2: <http://www.xbrl.org/2003/role/exampleRef> -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 49 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482785/280-10-55-49>Reference 3: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 270 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (i) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482964/270-10-50-1>Reference 4: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (ee) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482810/280-10-50-32>Reference 5: <http://fasb.org/us-gaap/role/ref/otherTransitionRef> -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (d) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482810/280-10-50-32>Reference 6: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482810/280-10-50-22>Reference 7: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 810 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph (bb) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481203/810-10-50-3>Reference 8: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 810 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 25 -Subparagraph (a) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481231/810-10-45-25>Reference 9: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 235 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.4-08(g)(1)(iii)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480678/235-10-S99-1>Reference 10: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 323 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph (c) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481687/323-10-50-3>Reference 11: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 825 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 28 -Subparagraph (f) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482907/825-10-50-28>Reference 12: <http://www.xbrl.org/2003/role/exampleRef> -Topic 852 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 10 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481372/852-10-55-10>Reference 13: <http://www.xbrl.org/2003/role/exampleRef> -Topic 946 -SubTopic 830 -Name Accounting Standards Codification -Section 55 -Paragraph 12 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479168/946-830-55-12>Reference 14: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(12)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478777/944-210-S99-1>Reference 15: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.6-04(8)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479170/946-210-S99-1>Reference 16: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(18)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1>Reference 17: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(4)(i)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1A>Reference 18: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(4)(iii)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1A>Reference 19: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(4)(iii)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1A>Reference 20: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(4)(iii)(A)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1A>Reference 21: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(4)(iv)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1A>Reference 22: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(5)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1A>Reference 23: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1B -Subparagraph (SX 210.13-02(a)(4)(i)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1B>Reference 24: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1B -Subparagraph (SX 210.13-02(a)(4)(iii)(A)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1B>Reference 25:

<https://asc.fasb.org/1943274/2147481404/852-10-50-7>Reference 29: <http://www.xbrl.org/2003/role/exampleRef> -Topic 880 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 30 -Subparagraph (d) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482810/280-10-50-30> + Details Name: us-gaap Liabilities Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionAmount of liabilities and equity items, including the portion of equity attributable to noncontrolling interests, if any. + ReferencesReference 1: <http://www.xbrl.org/2003/role/exampleRef> -Topic 852 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 10 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481372/852-10-55-10>Reference 2: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(25)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478777/944-210-S99-1>Reference 3: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 235 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.4-08(g)(1)(ii)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480678/235-10-S99-1>Reference 4: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 323 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph (c) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481687/323-10-50-3>Reference 5: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 825 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 28 -Subparagraph (f) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482907/825-10-50-28>Reference 6: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 942 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.9-03(23)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478546/942-210-S99-1>Reference 7: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(32)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1> + Details Name: us-gaap_LiabilitiesAndStockholdersEquity Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionTotal obligations incurred as part of normal operations that are expected to be paid during the following twelve months or within one business cycle, if longer. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(21)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1>Reference 2: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 810 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph (a) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481203/810-10-50-3>Reference 3: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 810 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 25 -Subparagraph (b) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481231/810-10-45-25>Reference 4: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 810 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph (bb) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481203/810-10-50-3>Reference 5: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 235 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.4-08(g)(1)(ii)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480678/235-10-S99-1>Reference 6: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 323 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph (c) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481687/323-10-50-3>Reference 7: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 825 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 28 -Subparagraph (f) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482907/825-10-50-28>Reference 8: <http://www.xbrl.org/2003/role/exampleRef> -Topic 852 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 10 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481372/852-10-55-10>Reference 9: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 5 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147483467/210-10-45-5>Reference 10: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(4)(i)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1>Reference 11: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(4)(ii)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1>Reference 12: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(4)(iii)(A)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1>Reference 13: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(4)(iv)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1>Reference 14: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(5)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1>Reference 15: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1B -Subparagraph (SX 210.13-02(a)(4)(i)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1>Reference 16: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1B -Subparagraph (SX 210.13-02(a)(4)(iii)(A)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1>Reference 17: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1B -Subparagraph (SX 210.13-02(a)(4)(iii)(B)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1>Reference 18: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1B -Subparagraph (SX 210.13-02(a)(4)(iv)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1>Reference 19: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1B -Subparagraph (SX 210.13-02(a)(5)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1>Reference 20: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 852 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 7 -Subparagraph (a) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481404/852-10-50-7>Reference 21: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 852 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 7 -Subparagraph (b) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481404/852-10-50-7> + Details Name: us-gaap LiabilitiesCurrent Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - ReferencesNo definition available. + Details Name: us-gaap LiabilitiesCurrentAbstract Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionAmount of investment in marketable security, classified as current. + ReferencesReference 1: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(2)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1> + Details Name: us-gaap MarketableSecuritiesCurrent Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionPresent value of lessee's discounted obligation for lease payments from operating lease, classified as current. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 842 -SubTopic 20 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Subparagraph (b) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479041/842-20-45-1> + Details Name: us-gaap OperatingLeaseLiabilityCurrent Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionPresent value of lessee's discounted obligation for lease payments from operating lease, classified as noncurrent. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 842 -SubTopic 20 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Subparagraph (b) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479041/842-20-45-1> + Details Name: us-gaap OperatingLeaseLiabilityNoncurrent Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionAmount of lessee's right to use underlying asset under operating lease. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 842 -SubTopic 20 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Subparagraph (a) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479041/842-20-45-1> + Details Name: us-gaap OperatingLeaseRightOfUseAsset Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionAmount of noncurrent assets classified as other. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(17)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1> + Details Name: us-gaap OtherAssetsNoncurrent Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionAggregate par or stated value of issued nonredeemable preferred stock (or preferred stock redeemable solely at the option of the issuer). This item includes treasury stock repurchased by the entity. Note: elements for number of nonredeemable preferred shares, par value and other disclosure concepts are in another section within stockholders' equity. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(28)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1>Reference 2: <http://www.xbrl.org/2003/role/exampleRef> -Topic 852 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 10 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481372/852-10-55-10>Reference 3: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(21)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478777/944-210-S99-1> + Details Name: us-gaap PreferredStockValue Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionAmount of asset related to consideration paid in advance for costs that provide economic benefits in future periods, and amount of other assets that are expected to be realized or consumed within one year or the normal operating cycle, if longer. + ReferencesReference 1: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(9)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1> + Details Name: us-gaap PrepaidExpenseAndOtherAssetsCurrent Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionAmount after accumulated depreciation, depletion and amortization of physical assets used in the normal conduct of business to produce goods and services and not intended for resale. Examples include, but are not limited to, land, buildings, machinery and equipment, office equipment, and furniture and fixtures. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 50 -Paragraph 1 -SubTopic 10 -Topic 360 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482099/360-10-50-1>Reference 2: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 842 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 7A -Subparagraph (a) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478964/842-20-50-7>Reference 3: <http://www.xbrl.org/2003/role/exampleRef> -Topic 852 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 10 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481372/852-10-55-10>Reference 4: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(8)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478777/944-210-S99-1>Reference 5: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 942 -SubTopic 360 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478451/942-360-50-1> + Details Name: us-gaap PropertyPlantAndEquipmentNet Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionAmount of accumulated undistributed earnings (deficit). + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(30)(a)(3)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1>Reference 2: <http://www.xbrl.org/2003/role/exampleRef> -Topic 852 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 10 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481372/852-10-55-10>Reference 3: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 944 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 2 -Subparagraph (g)(2)(i) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480016/944-40-65-2>Reference 4: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 944 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 2 -Subparagraph (h)(2) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480016/944-40-65-2>Reference 5: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 946 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 11 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480990/946-20-50-11>Reference 6: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(23)(a)(4)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478777/944-210-S99-1>Reference 7: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.6-04(17)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479170/946-210-S99-1>Reference 8: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.3-04) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480008/505-10-S99-1> + Details Name: us-gaap RetainedEarningsAccumulatedDeficit Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionReflects the total carrying amount as of the balance sheet date of debt having initial terms less than one year or the normal operating cycle, if longer. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(19)(a)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1>Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 942 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.9-03(13)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478546/942-210-S99-1>Reference 3: <http://www.xbrl.org/2003/role/exampleRef> -Topic 852 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 10 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481372/852-10-55-10>Reference 4: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(16)(a)(1)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478777/944-210-S99-1> + Details Name: us-gaap_ShortTermBorrowings Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionAmount of equity (deficit) attributable to parent. Excludes temporary equity and equity attributable to noncontrolling interest. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(29)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1>Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(30)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479170/946-210-S99-2>Reference 3: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-09(6)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479134/946-220-S99-3>Reference 10: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-09(7)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479134/946-220-S99-3>Reference 11: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 235 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.4-08(g)(1)(ii)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480678/235-10-S99-1>Reference 12: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 323 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph (c) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481687/323-10-50-3>Reference 13: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 825 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 28 -Subparagraph (f) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482907/825-10-50-28>Reference 14: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 310 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -

Paragraph 2 - Subparagraph (SAB Topic 4.E) - Publisher FASB -URI https://asc.fasb.org/1943274/2147480418/310-10-S99-2 + Details Name: us-gaap StockholdersEquityNamespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - ReferencesNo definition available. + Details Name: us-gaap StockholdersEquityAbstractNamespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration XML 10 R3.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Consolidated Balance Sheets (Parenthetical) (Unaudited) - USD (\$) in in Thousands Jun. 30, 2024 Dec. 31, 2023 Accounts Receivable, Allowance for Credit Loss \$ 0 Common stock and additional paid in capital, par value (in Dollars per share) \$ 0.0001 \$ 0.0001 Common stock and additional paid in capital, shares issued 248,624,729 219,046,219 Common stock and additional paid in capital, shares outstanding 248,624,729 219,046,219 Property and equipment, net of accumulated depreciation of \$782 and \$666, respectively \$ 782 \$ 666 Intangible assets, net of amortization of \$138 \$ 138 A Additional Paid-in Capital [Member] A A Common stock and additional paid in capital, par value (in Dollars per share) \$ 0.0001 \$ 0.0001 Common stock and additional paid in capital, shares authorized 1,060,000,000 1,060,000,000 Class A Common Stock [Member] A A Common stock and additional paid in capital, shares issued 247,624,729 218,046,219 Common stock and additional paid in capital, shares outstanding 247,624,729 218,046,219 Class A Common Stock [Member] | Additional Paid-in Capital [Member] A A Common stock and additional paid in capital, shares authorized 1,000,000,000 1,000,000,000 Class B Common Stock [Member] A A Common stock and additional paid in capital, shares issued 1,000,000 1,000,000 Common stock and additional paid in capital, shares outstanding 1,000,000 1,000,000 Class B Common Stock [Member] | Additional Paid-in Capital [Member] A A Common stock and additional paid in capital, shares authorized 60,000,000 60,000,000 Convertible Preferred Stock [Member] A A Preferred stock par value \$ 0.0001 \$ 0.0001 Preferred Stock, Shares Issued 851,202 851,202 Preferred Stock Shares Outstanding 851,202 851,202 X - DefinitionIntangible Assets Net of Amortization + ReferencesNo definition available. + Details Name: nuvb IntangibleAssetsNetOfAmortization Namespace Prefix: nuvb Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionAmount of accumulated depreciation, depletion and amortization for physical assets used in the normal conduct of business to produce goods and services. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(8)(b)) -Publisher FASB -URI https://asc.fasb.org/1943274/214748777/944-210-S99-1Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(14)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 3: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Topic 360 -SubTopic 10 -Section 50 -Paragraph 1 -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482099/360-10-50-1 + Details Name: us-gaap AccumulatedDepreciationDepletionAndAmortizationPropertyPlantAndEquipmentNamespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionAmount of allowance for credit loss on accounts receivable. + ReferencesReference 1: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(5)) -Publisher FASB -URI https://asc.fasb.org/1943274/214748777/944-210-S99-1Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(4)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 326 -SubTopic 20 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479344/326-20-45-1Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Topic 310 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481962/310-10-50-4Reference 5: http://www.xbrl.org/2003/role/disclosureRef -Topic 326 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 13 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479319/326-20-50-13Reference 6: http://www.xbrl.org/2003/role/disclosureRef -Topic 326 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 13 -Subparagraph (f) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479319/326-20-50-13 + Details Name: us-gaap AllowanceForDoubtfulAccountsReceivableNamespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionFace amount or stated value per share of common stock. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(29)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1 + Details Name: us-gaap CommonStockParOrStatedValuePerShareNamespace Prefix: us-gaap Data Type: dtr-types:perShareItemType Balance Type: na Period Type: instant X - DefinitionThe maximum number of common shares permitted to be issued by an entity's charter and bylaws. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(29)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.6-04(16)(a)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479170/946-210-S99-1 + Details Name: us-gaap CommonStockSharesAuthorizedNamespace Prefix: us-gaap Data Type: xbrli:sharesItemType Balance Type: na Period Type: instant X - DefinitionTotal number of common shares of an entity that have been sold or granted to shareholders (includes common shares that were issued, repurchased and remain in the treasury). These shares represent capital invested by the firm's shareholders and owners, and may be all or only a portion of the number of shares authorized. Shares issued include shares outstanding and shares held in the treasury. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(29)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.6-04(16)(a)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-3Reference 5: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.6-04(16)(a)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479170/946-210-S99-1Reference 6: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-09(4)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-3 + Details Name: us-gaap CommonStockSharesOutstandingNamespace Prefix: us-gaap Data Type: xbrli:sharesItemType Balance Type: na Period Type: instant X - DefinitionFace amount or stated value per share of preferred stock nonredeemable or redeemable solely at the option of the issuer. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(28)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 13 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481112/505-10-50-2Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(29)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph (SX 210.6-05(4)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479170/946-210-S99-2Reference 4: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-09(4)(b)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-3Reference 5: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.6-04(16)(a)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479170/946-210-S99-1Reference 6: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-09(4)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-3 + Details Name: us-gaap PreferredStockParOrStatedValuePerShareNamespace Prefix: us-gaap Data Type: dtr-types:perShareItemType Balance Type: na Period Type: instant X - DefinitionNumber of shares issued for nonredeemable preferred shares and preferred shares redeemable solely at option of issuer. Includes, but is not limited to, preferred shares issued, repurchased, and held as treasury shares. Excludes preferred shares classified as debt. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(28)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 13 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481112/505-10-50-13 + Details Name: us-gaap PreferredStockSharesIssuedNamespace Prefix: us-gaap Data Type: xbrli:sharesItemType Balance Type: na Period Type: instant X - DefinitionAggregate share number for all nonredeemable preferred stock (or preferred stock redeemable solely at the option of the issuer) held by stockholders. Does not include preferred shares that have been repurchased. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(28)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph (SX 210.6-05(4)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479170/946-210-S99-2Reference 3: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-09(4)(b)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-3Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.6-04(16)(a)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-3 + Details Name: us-gaap PreferredStockSharesOutstandingNamespace Prefix: us-gaap Data Type: xbrli:sharesItemType Balance Type: na Period Type: instant X - Details Name: us-gaap StatementEquityComponentsAxis=us-gaap AdditionalPaidInCapitalMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap StatementClassOfStockAxis=us-gaap CommonClassAMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap StatementClassOfStockAxis=us-gaap ConvertiblePreferredStockMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 11 R4.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Consolidated Statements of Operations and Comprehensive Loss (Unaudited) - USD (\$) in in Thousands 3 Months Ended 6 Months Ended Jun. 30, 2024 Jun. 30, 2023 Jun. 30, 2024 Jun. 30, 2023 Income Statement [Abstract] A A A Revenue \$ 1,435 \$ 0 \$ 1,435 \$ 0 Cost of revenue 1,347 A 1,347 A Gross Profit 88 0 88 0 Operating expenses: A A A Research and development 29,247 18,590 42,089 37,377 Acquired in-process research and development 425,070 0 425,070 0 Selling, general and administrative 16,156 7,541 23,513 15,275 Total operating expenses 47,047 26,131 490,672 A Loss from operations (470,385) (26,131) (490,584) (52,652) Other income (expense): A A A Interest income 7,144 6,086 14,274 11,065 Interest expense (132) 0 (132) 0 Investment advisory fees (247) (231) (512) (461) Change in fair value of warrant liability 1,135 (265) (324) (123) Realized loss on marketable securities (7) (99) (6) (195) Total other income (expense), net 7,893 5,491 13,300 10,286 Loss before income taxes (462,492) (20,640) (477,284) (42,366) Provision for income taxes 0 0 0 0 Net loss \$ (462,492) \$ (20,640) \$ (477,284) \$ (42,366) Net loss per share attributable to common stockholders, basic \$ (1.89) \$ (0.09) \$ (2.06) \$ (0.19) Net loss per share attributable to common stockholders, diluted \$ (1.89) \$ (0.09) \$ (2.06) \$ (0.19) Weighted average common shares outstanding, basic 244,738 218,848 231,893 218,795 Weighted average common shares outstanding, diluted 244,738 218,848 231,893 218,795 Comprehensive loss: A A A Net loss \$ (462,492) \$ (20,640) \$ (477,284) \$ (42,366) Other comprehensive loss, net of taxes: A A A Currency translation adjustment 148 0 148 0 Change in unrealized (loss) gain on available-for-sale securities (245) (1,457) (1,595) 1,131 Comprehensive loss \$ (462,589) \$ (22,079) \$ (478,731) \$ (41,235) X - DefinitionAcquired in-process research and development + ReferencesNo definition available. + Details Name: nuvb AcquiredInProcessResearchAndDevelopmentNamespace Prefix: nuvb Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionAmount after tax of increase (decrease) in equity from transactions and other events and circumstances from net income and other comprehensive income, attributable to parent entity. Excludes changes in equity resulting from investments by owners and distributions to owners. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph (SX 210.5-03(24)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-S99-2Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 942 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.9-04(26)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478524/942-220-S99-1Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-04(22)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477250/944-220-S99-1Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 1A -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482790/220-10-45-1AReference 5: http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 1B -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482790/220-10-45-1BReference 6: http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-04(20)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477250/944-220-S99-1Reference 7: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Topic 220 -SubTopic 10 -Section 45 -Paragraph 5 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482790/220-10-45-5 + Details Name: us-gaap ComprehensiveIncomeNetOfTaxNamespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - ReferencesNo definition available. + Details Name: us-gaap ComprehensiveIncomeNetOfTaxAbstractNamespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionThe aggregate costs related to goods produced and sold and services rendered by an entity during the reporting period. This excludes costs incurred during the reporting period related to financial services rendered and other revenue generating activities. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph (SX 210.5-03(2)(d)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-S99-2Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph (SX 210.5-03(2)(a)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-S99-2Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 924 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SAB Topic 11.L) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479941/924-10-S99-1 + Details Name: us-gaap CostOfGoodsAndServicesSoldNamespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionAmount of unrealized gain (loss) on investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available-for-sale), investment in debt security measured at amortized cost (held-to-maturity) and investment in debt security measured at fair value with change in fair value recognized in net income (trading). + ReferencesReference 1: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph (SX 210.5-03(9)(a)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-S99-2Reference 2: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph (SX 210.5-03(7)(c)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-S99-2Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-04(3)(d)(2)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477250/944-220-S99-1 + Details Name: us-gaap DebtSecuritiesUnrealizedGainLossNamespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionThe amount of net income (loss) for the period per each share of common stock or unit outstanding during the reporting period. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 6 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-6Reference 2: http://www.xbrl.org/2003/role/exampleRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 52 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482635/260-10-55-52Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 805 -SubTopic 60 -Name Accounting Standards Codification -Section 65 -Paragraph 1 -Subparagraph (g) -Publisher FASB -URI https://asc.fasb.org/1943274/2147476176/805-60-65-1Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Topic 740 -SubTopic 323 -Name Accounting Standards Codification -Section 65 -Paragraph 2 -Subparagraph (g)(3) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478666/740-323-65-2Reference 5: http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -

Paragraph 3 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147483443/250-10-50-3Reference> 6: <http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 15 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482635/260-10-55-15Reference> 7: [http://www.xbrl.org/2003/role/disclosureRef -Topic 815 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 1 -Subparagraph \(e\)\(4\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480175/815-40-65-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 815 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 1 -Subparagraph (e)(4) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480175/815-40-65-1Reference) 8: [http://www.xbrl.org/2003/role/disclosureRef -Topic 815 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 1 -Subparagraph \(f\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480175/815-40-65-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 815 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 1 -Subparagraph (f) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480175/815-40-65-1Reference) 9: [http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 11 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-11Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 11 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-11Reference) 10: [http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 11 -Subparagraph \(b\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-11Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 11 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-11Reference) 11: [http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 7 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-7Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 7 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-7Reference) 12: <http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 2 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482689/260-10-45-2Reference> 13: [http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 60B -Subparagraph \(d\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482689/260-10-45-60BReference](http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 60B -Subparagraph (d) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482689/260-10-45-60BReference) 14: <http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-4Reference> 15: [http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482662/260-10-50-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482662/260-10-50-1Reference) 16: <http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 10 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482689/260-10-45-10Reference> 17: [http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph \(SX 210.5-03\(25\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-S99-2Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph (SX 210.5-03(25)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-S99-2Reference) 18: [http://www.xbrl.org/2003/role/disclosureRef -Topic 942 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.9-04\(27\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478524/942-220-S99-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 942 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.9-04(27)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478524/942-220-S99-1Reference) 19: [http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.7-04\(23\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477250/944-220-S99-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-04(23)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477250/944-220-S99-1Reference) 20: <http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 7 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482689/260-10-45-7> + Details Name: us-gaap_EarningsPerShareBasic Namespace Prefix: us-gaap_Data Type: dtr-types:perShareItemType Balance Type: na Period Type: duration X - DefinitionThe amount of net income (loss) for the period available to each share of common stock or common unit outstanding during the reporting period and to each share or unit that would have been outstanding assuming the issuance of common shares or units for all dilutive potential common shares or units outstanding during the reporting period. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 6 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-6Reference> 2: <http://www.xbrl.org/2003/role/exampleRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 52 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482635/260-10-55-52Reference> 3: [http://www.xbrl.org/2003/role/disclosureRef -Topic 805 -SubTopic 60 -Name Accounting Standards Codification -Section 65 -Paragraph 1 -Subparagraph \(g\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147476176/805-60-65-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 805 -SubTopic 60 -Name Accounting Standards Codification -Section 65 -Paragraph 1 -Subparagraph (g) -Publisher FASB -URI https://asc.fasb.org/1943274/2147476176/805-60-65-1Reference) 4: [http://www.xbrl.org/2003/role/disclosureRef -Topic 740 -SubTopic 323 -Name Accounting Standards Codification -Section 65 -Paragraph 2 -Subparagraph \(g\)\(3\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478666/740-323-65-2Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 740 -SubTopic 323 -Name Accounting Standards Codification -Section 65 -Paragraph 2 -Subparagraph (g)(3) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478666/740-323-65-2Reference) 5: <http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-3Reference> 6: <http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 15 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482635/260-10-55-15Reference> 7: [http://www.xbrl.org/2003/role/disclosureRef -Topic 815 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 1 -Subparagraph \(e\)\(4\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480175/815-40-65-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 815 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 1 -Subparagraph (e)(4) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480175/815-40-65-1Reference) 8: [http://www.xbrl.org/2003/role/disclosureRef -Topic 815 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 1 -Subparagraph \(f\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480175/815-40-65-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 815 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 1 -Subparagraph (f) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480175/815-40-65-1Reference) 9: [http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 11 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-11Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 11 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-11Reference) 10: [http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 7 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-7Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 7 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-7Reference) 12: <http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 2 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482689/260-10-45-2Reference> 13: [http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 60B -Subparagraph \(d\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482689/260-10-45-60BReference](http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 60B -Subparagraph (d) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482689/260-10-45-60BReference) 14: <http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-4Reference> 15: [http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482662/260-10-50-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482662/260-10-50-1Reference) 16: [http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph \(SX 210.5-03\(25\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-S99-2Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph (SX 210.5-03(25)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-S99-2Reference) 17: [http://www.xbrl.org/2003/role/disclosureRef -Topic 942 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.9-04\(27\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478524/942-220-S99-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 942 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.9-04(27)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478524/942-220-S99-1Reference) 18: [http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.7-04\(23\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477250/944-220-S99-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-04(23)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477250/944-220-S99-1Reference) 19: <http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 7 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482689/260-10-45-7> + Details Name: us-gaap_EarningsPerShareDiluted Namespace Prefix: us-gaap_Data Type: dtr-types:perShareItemType Balance Type: na Period Type: duration X - DefinitionAmount of expense (income) related to adjustment to fair value of warrant liability. + ReferencesReference 1: [http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Subparagraph \(b\) -SubTopic 10 -Topic 230 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482740/230-10-45-28Reference](http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Subparagraph (b) -SubTopic 10 -Topic 230 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482740/230-10-45-28Reference) 2: <http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Section 25 -Paragraph 13 -SubTopic 10 -Topic 480 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481766/480-10-25-13> + Details Name: us-gaap_FairValueAdjustmentOfWarrants Namespace Prefix: us-gaap_Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionAggregate revenue less cost of goods and services sold or operating expenses directly attributable to the revenue generation activity. + ReferencesReference 1: <http://www.xbrl.org/2009/role/commonPracticeRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22Reference> 2: <http://www.xbrl.org/2003/role/exampleRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 48 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482785/280-10-55-48Reference> 3: [http://www.xbrl.org/2009/role/commonPracticeRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph \(f\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference](http://www.xbrl.org/2009/role/commonPracticeRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (f) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference) 4: [http://www.xbrl.org/2009/role/commonPracticeRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 30 -Subparagraph \(b\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482964/270-10-50-30Reference](http://www.xbrl.org/2009/role/commonPracticeRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 30 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482964/270-10-50-30Reference) 5: [http://www.xbrl.org/2009/role/commonPracticeRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph \(ee\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference](http://www.xbrl.org/2009/role/commonPracticeRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (ee) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference) 7: [http://fasb.org/us-gaap/role/ref/otherTransitionRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph \(c\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference](http://fasb.org/us-gaap/role/ref/otherTransitionRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference) 8: [http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.7-04\(11\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477250/944-220-S99-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-04(11)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477250/944-220-S99-1Reference) 9: <http://www.xbrl.org/2003/role/exampleRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 31 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-31Reference> 10: [http://www.xbrl.org/2003/role/disclosureRef -Topic 942 -SubTopic 235 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.9-05\(b\)\(2\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477314/942-235-S99-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 942 -SubTopic 235 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.9-05(b)(2)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477314/942-235-S99-1Reference) 11: [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph \(SX 210.5-03\(10\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-S99-2Reference](http://fasb.org/us-gaap/role/ref/legacyRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph (SX 210.5-03(10)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-S99-2Reference) 12: [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 942 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.9-04\(15\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478524/942-220-S99-1](http://fasb.org/us-gaap/role/ref/legacyRef -Topic 942 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.9-04(15)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478524/942-220-S99-1) + Details Name: us-gaap_IncomeLossFromContinuingOperationsBeforeIncomeTaxesExtraordinaryItemsNoncontrollingInterest Namespace Prefix: us-gaap_Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - ReferencesNo definition available. + Details Name: us-gaap_IncomeStatementAbstract Namespace Prefix: us-gaap_Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionAmount of current income tax expense (benefit) and deferred income tax expense (benefit) pertaining to continuing operations. + ReferencesReference 1: [http://www.xbrl.org/2003/role/disclosureRef -Topic 270 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph \(i\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482964/270-10-50-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 270 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (i) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482964/270-10-50-1Reference) 2: [http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph \(g\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (g) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference) 3: <http://www.xbrl.org/2003/role/disclosureRef -Topic 740 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 12 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482685/740-10-50-12Reference> 4: <http://www.xbrl.org/2003/role/exampleRef -Topic 740 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 231 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482663/740-10-55-231Reference> 5: <http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 9 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-9Reference> 6: [http://www.xbrl.org/2003/role/disclosureRef -Topic 740 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SAB Topic 6.1.7\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479360/740-10-S99-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 740 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SAB Topic 6.1.7) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479360/740-10-S99-1Reference) 7: <http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 8 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-8Reference> 8: <http://www.xbrl.org/2003/role/disclosureRef -Topic 740 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 10 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482685/740-10-50-10Reference> 9: [http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph \(h\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph (h) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22Reference)

22Reference 10: [http://www.xbrl.org/2003/role/disclosureRef-Topic 944-SubTopic 220-Name Accounting Standards Codification-Section S99-Paragraph 1-Subparagraph \(SX 210.7-04\(9\)\)](http://www.xbrl.org/2003/role/disclosureRef-Topic 944-SubTopic 220-Name Accounting Standards Codification-Section S99-Paragraph 1-Subparagraph (SX 210.7-04(9))) - Publisher FASB -URI <https://asc.fasb.org/1943274/2147477250/944-220-S99-1Reference 11>: [http://www.xbrl.org/2003/role/disclosureRef-Name Accounting Standards Codification-Section 45-Paragraph 2-Subparagraph \(a\)-SubTopic 20-Topic 740-Publisher FASB-URI https://asc.fasb.org/1943274/2147482659/740-20-45-2Reference 12](http://www.xbrl.org/2003/role/disclosureRef-Name Accounting Standards Codification-Section 45-Paragraph 2-Subparagraph (a)-SubTopic 20-Topic 740-Publisher FASB-URI https://asc.fasb.org/1943274/2147482659/740-20-45-2Reference 12):

codification -Section 50 -Paragraph 15 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-50Reference 8: [http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 30 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-30](http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 30 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-30)Reference 9: <http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 42 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-42>Reference 10: [http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph \(b\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22](http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22)Reference 11: [http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22](http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22)Reference 12: [http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22](http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22)Reference 13: [http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-4](http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-4) + Details Name: us-gaap RevenueFromContractWithCustomerExcludingAssessedTax Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionThe aggregate total costs related to selling a firm's product and services, as well as all other general and administrative expenses. Direct selling expenses (for example, credit, warranty, and advertising) are expenses that can be directly linked to the sale of specific products. Indirect selling expenses are expenses that cannot be directly linked to the sale of specific products, for example telephone expenses, Internet, and postal charges. General and administrative expenses include salaries of non-sales personnel, rent, utilities, communication, etc. + ReferencesReference 1: [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph \(SX 210.5-03\(4\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-50-39-2](http://fasb.org/us-gaap/role/ref/legacyRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph (SX 210.5-03(4)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-50-39-2) + Details Name: us-gaap SellingGeneralAndAdministrativeExpense Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionAmount of income tax expense (benefit) from translating foreign currency financial statements into the reporting currency. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Section 45 -Paragraph 12 -SubTopic 10 -Topic 220 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482790/220-10-45-12>Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 11 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482790/220-10-45-11>Reference 3: [http://www.xbrl.org/2009/role/commonPracticeRef -Topic 830 -SubTopic 30 -Name Accounting Standards Codification -Section 45 -Paragraph 20 -Subparagraph \(c\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481694/830-30-45-20](http://www.xbrl.org/2009/role/commonPracticeRef -Topic 830 -SubTopic 30 -Name Accounting Standards Codification -Section 45 -Paragraph 20 -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481694/830-30-45-20) + Details Name: us-gaap TranslationAdjustmentFunctionalToReportingCurrencyTaxExpenseBenefit Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionThe average number of shares or units issued and outstanding that are used in calculating diluted EPS or earnings per unit (EPU), determined based on the timing of issuance of shares or units in the period. + ReferencesReference 1: [http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482662/260-10-50-1](http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482662/260-10-50-1)Reference 2: <http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 10 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482689/260-10-45-10> + Details Name: us-gaap WeightedAverageNumberOfDilutedSharesOutstandingBasic Namespace Prefix: us-gaap Data Type: xbrli:sharesItemType Balance Type: na Period Type: duration X - DefinitionNumber of (basic) shares or units, after adjustment for contingently issuable shares or units and other shares or units not deemed outstanding, determined by relating the portion of time within a reporting period that common shares or units have been outstanding to the total time in that period. + ReferencesReference 1: [http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482662/260-10-50-1](http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482662/260-10-50-1)Reference 2: <http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 10 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482689/260-10-45-10> + Details Name: us-gaap WeightedAverageNumberOfSharesOutstandingBasic Namespace Prefix: us-gaap Data Type: xbrli:sharesItemType Balance Type: na Period Type: duration XML 12 R5.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Consolidated Statements of Redeemable Convertible Preferred Stock and Stockholders' Equity (Unaudited) - USD (\$) \$ in Thousands Total Preferred Class A [Member] Common Stock Including Additional Paid In Capital [Member] Common Stock Including Additional Paid In Capital [Member] Class A Shares [Member] Common Stock Including Additional Paid In Capital [Member] Class B Shares [Member] Convertible Preferred Stocks [Member] Preferred Class A [Member] Accumulated Deficit [Member] Accumulated Other Comprehensive Income [Member] Balance at Dec. 31, 2022 \$ 655,076 \$ 927,604 \$ 0 \$ (267,002) \$ (5,526) Balance (in Shares) at Dec. 31, 2022 \$ 4,837 \$ 217,632,699 1,000,000 \$ 0 \$ Exercise of stock options, shares \$ 0 \$ 171,023 \$ 0 \$ Exercise of stock options 298 \$ 298 \$ 0 \$ Stock-based compensation 4,837 \$ 4,837 \$ 0 \$ Net Income (Loss) (21,726) \$ 0 \$ (21,726) Other comprehensive loss 2,588 \$ 0 \$ 2,588 Balance at Mar. 31, 2023 641,073 \$ 932,739 \$ 0 \$ (288,728) (2,938) Balance (in Shares) at Mar. 31, 2023 \$ 4,837 \$ 217,632,699 1,000,000 \$ 0 \$ Currency Translation Adjustment 0 \$ 0 \$ 0 \$ Net Income (Loss) (42,366) \$ 0 \$ 0 \$ Balance at Jun. 30, 2023 624,632 \$ 938,395 \$ 0 \$ (309,368) (4,395) Balance (in Shares) at Jun. 30, 2023 \$ 4,837 \$ 217,899,462 1,000,000 \$ 0 \$ Balance at Mar. 31, 2023 641,073 \$ 932,739 \$ 0 \$ (288,728) (2,938) Balance (in Shares) at Mar. 31, 2023 \$ 4,837 \$ 217,899,462 1,000,000 \$ 0 \$ Issuance of common stock for purchase under ESPP, (shares) \$ 0 \$ 95,740 \$ 0 \$ Issuance of common stock for purchase under ESPP 133 \$ 133 \$ 0 \$ Stock-based compensation 5,523 \$ 5,523 \$ 0 \$ Net Income (Loss) (20,640) \$ 0 \$ (20,64

[http://www.xbrl.org/2003/role/disclosureRef-Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1B -Subparagraph \(SX 210.13-02\(a\)\(5\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480097/470-10-S99-1BReference 32:](http://www.xbrl.org/2003/role/disclosureRef-Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1B -Subparagraph (SX 210.13-02(a)(5)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480097/470-10-S99-1BReference 32:) [http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 60B -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482689/260-10-45-60BReference 33:](http://www.xbrl.org/2003/role/disclosureRef -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 60B -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482689/260-10-45-60BReference 33:) <http://www.xbrl.org/2003/role/disclosureRef -Topic 205 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 7 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483499/205-20-50-7Reference 34:> <http://fasb.org/us-gaap/role/ref/legacyRef -Topic 230 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482740/230-10-45-28Reference 35:> [http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 1A -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482790/220-10-45-1AReference 36:](http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 1A -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482790/220-10-45-1AReference 36:) [http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 1B -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482790/220-10-45-1BReference 37:](http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 1B -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482790/220-10-45-1BReference 37:) [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 942 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.9-04\(22\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478524/942-220-S99-1 + Details Name: us-gaap_NetIncomeLoss Namespace Prefix: us-gaap_Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionAmount after tax and reclassification adjustments of other comprehensive income \(loss\). + ReferencesReference 1:](http://fasb.org/us-gaap/role/ref/legacyRef -Topic 942 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.9-04(22)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478524/942-220-S99-1 + Details Name: us-gaap_NetIncomeLoss Namespace Prefix: us-gaap_Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionAmount after tax and reclassification adjustments of other comprehensive income (loss). + ReferencesReference 1:) <http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482765/220-10-50-4Reference 7:> <http://www.xbrl.org/2009/role/commonPracticeRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 5 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482765/220-10-50-5Reference 8:> <http://www.xbrl.org/2003/role/disclosureRef -Topic 830 -SubTopic 30 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481674/830-30-45-1Reference 4:> <http://www.xbrl.org/2003/role/disclosureRef -Topic 830 -SubTopic 30 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481694/830-30-45-17Reference 5:> [http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.7-04\(19\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477250/944-220-S99-1Reference 6:](http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-04(19)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477250/944-220-S99-1Reference 6:) <http://www.xbrl.org/2009/role/commonPracticeRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482765/220-10-50-4Reference 7:> <http://www.xbrl.org/2009/role/commonPracticeRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 5 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482765/220-10-50-5Reference 8:> <http://www.xbrl.org/2003/role/disclosureRef -Topic 830 -SubTopic 30 -Name Accounting Standards Codification -Section 45 -Paragraph 20 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481694/830-30-45-20Reference 9:> [http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph \(SX 210.5-03\(21\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-S99-2Reference 10:](http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph (SX 210.5-03(21)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-S99-2Reference 10:) [http://www.xbrl.org/2003/role/disclosureRef -Topic 942 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.9-04\(23\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478524/942-220-S99-1Reference 11:](http://www.xbrl.org/2003/role/disclosureRef -Topic 942 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.9-04(23)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478524/942-220-S99-1Reference 11:) [http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 1B -Subparagraph \(b\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482790/220-10-45-1BReference 12:](http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 1B -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482790/220-10-45-1BReference 12:) <http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Section 55 -Paragraph 15 -SubTopic 10 -Topic 220 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482739/220-10-55-15Reference 13:> [http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 1A -Subparagraph \(b\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482790/220-10-45-1A + Details Name: us-gaap_OtherComprehensiveIncomeLossNetOfTax Namespace Prefix: us-gaap_Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionNumber of shares issued which are neither cancelled nor held in the treasury. + ReferencesNo definition available. + Details Name: us-gaap_SharesOutstanding Namespace Prefix: us-gaap_Data Type: xbrli:sharesItem Type Balance Type: na Period Type: instant X - DefinitionNumber of share options \(or share units\) exercised during the current period. + ReferencesReference 1:](http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 1A -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482790/220-10-45-1A + Details Name: us-gaap_OtherComprehensiveIncomeLossNetOfTax Namespace Prefix: us-gaap_Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionNumber of shares issued which are neither cancelled nor held in the treasury. + ReferencesNo definition available. + Details Name: us-gaap_SharesOutstanding Namespace Prefix: us-gaap_Data Type: xbrli:sharesItem Type Balance Type: na Period Type: instant X - DefinitionNumber of share options (or share units) exercised during the current period. + ReferencesReference 1:) <http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Section 50 -Paragraph 2 -SubTopic 10 -Topic 505 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481112/505-10-50-2Reference 2:> [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.5-02\(28\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 3:](http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(28)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 3:) [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.5-02\(29\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 4:](http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(29)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 4:) [http://www.xbrl.org/2003/role/disclosureRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph \(c\)\(1\)\(iv\)\(02\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2Reference 5:](http://www.xbrl.org/2003/role/disclosureRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(iv)(02) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2Reference 5:) [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.3-04\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480008/505-10-S99-1 + Details Name: us-gaap_StockIssuedDuringPeriodSharesStockOptionsExercised Namespace Prefix: us-gaap_Data Type: xbrli:sharesItem Type Balance Type: na Period Type: duration X - DefinitionValue of stock issued as a result of the exercise of stock options. + ReferencesReference 1:](http://fasb.org/us-gaap/role/ref/legacyRef -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.3-04) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480008/505-10-S99-1 + Details Name: us-gaap_StockIssuedDuringPeriodSharesStockOptionsExercised Namespace Prefix: us-gaap_Data Type: xbrli:sharesItem Type Balance Type: na Period Type: duration X - DefinitionValue of stock issued as a result of the exercise of stock options. + ReferencesReference 1:) <http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -SubTopic 10 -Topic 505 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481112/505-10-50-2Reference 2:> [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.5-02\(29\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 3:](http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(29)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 3:) [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.5-02\(30\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 4:](http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(30)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 4:) [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.5-02\(31\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 5:](http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(31)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 5:) [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.3-04\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480008/505-10-S99-1 + Details Name: us-gaap_StockIssuedDuringPeriodValueStockOptionsExercised Namespace Prefix: us-gaap_Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionAmount of equity \(deficit\) attributable to parent. Excludes temporary equity and equity attributable to noncontrolling interest. + ReferencesReference 1:](http://fasb.org/us-gaap/role/ref/legacyRef -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.3-04) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480008/505-10-S99-1 + Details Name: us-gaap_StockIssuedDuringPeriodValueStockOptionsExercised Namespace Prefix: us-gaap_Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionAmount of equity (deficit) attributable to parent. Excludes temporary equity and equity attributable to noncontrolling interest. + ReferencesReference 1:) [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.5-02\(29\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 2:](http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(29)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 2:) [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.5-02\(30\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 3:](http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(30)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 3:) [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.5-02\(31\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 4:](http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(31)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 4:) <http://www.xbrl.org/2003/role/exampleRef -Topic 852 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 10 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481372/852-10-55-10Reference 5:> <http://www.xbrl.org/2003/role/exampleRef -Topic 946 -SubTopic 830 -Name Accounting Standards Codification -Section 55 -Paragraph 12 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479168/946-830-55-12Reference 6:> [http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.6-04\(19\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479170/946-210-S99-1Reference 7:](http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.6-04(19)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479170/946-210-S99-1Reference 7:) [http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph \(SX 210.6-05\(4\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479170/946-210-S99-2Reference 8:](http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph (SX 210.6-05(4)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479170/946-210-S99-2Reference 8:) [http://www.xbrl.org/2009/role/commonPracticeRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph \(SX 210.6-09\(4\)\(b\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-3Reference 9:](http://www.xbrl.org/2009/role/commonPracticeRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-09(4)(b)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-3Reference 9:) [http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph \(SX 210.6-09\(6\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-3Reference 10:](http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-09(6)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-3Reference 10:) [http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph \(SX 210.6-09\(7\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-3Reference 11:](http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-09(7)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-3Reference 11:) [http://www.xbrl.org/2009/role/commonPracticeRef -Topic 235 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.4-08\(g\)\(1\)\(iii\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480678/235-10-S99-1Reference 12:](http://www.xbrl.org/2009/role/commonPracticeRef -Topic 235 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.4-08(g)(1)(iii)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480678/235-10-S99-1Reference 12:) [http://www.xbrl.org/2009/role/commonPracticeRef -Topic 323 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph \(c\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481687/323-10-50-3Reference 13:](http://www.xbrl.org/2009/role/commonPracticeRef -Topic 323 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481687/323-10-50-3Reference 13:) [http://www.xbrl.org/2009/role/commonPracticeRef -Topic 825 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 28 -Subparagraph \(f\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482907/825-10-50-28Reference 14:](http://www.xbrl.org/2009/role/commonPracticeRef -Topic 825 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 28 -Subparagraph (f) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482907/825-10-50-28Reference 14:)

45 -Paragraph 28 -Subparagraph (b) -SubTopic 10 -Topic 230 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-28>Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Topic 360 -SubTopic 10 -Section 50 -Paragraph 1 -Subparagraph (a) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482099/360-10-50-1> + Details Name: us-gaap DepreciationAndAmortization Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionAmount of increase (decrease) from effect of exchange rate changes on cash and cash equivalents, and cash and cash equivalents restricted to withdrawal or usage; held in foreign currencies. Excludes amounts for disposal group and discontinued operations. Cash includes, but is not limited to, currency on hand, demand deposits with banks or financial institutions, and other accounts with general characteristics of demand deposits. Cash equivalents include, but are not limited to, short-term, highly liquid investments that are both readily convertible to known amounts of cash and so near their maturity that they present insignificant risk of changes in value because of changes in interest rates. + ReferencesReference 1: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 830 -SubTopic 230 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147477401/830-230-45-1> + Details Name: us-gaap_EffectOfExchangeRateOnCashCashEquivalentsRestrictedCashAndRestrictedCashEquivalents Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionAmount of expense (income) related to adjustment to fair value of warrant liability. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Subparagraph (b) -SubTopic 10 -Topic 230 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-28>Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 25 -Paragraph 13 -SubTopic 10 -Topic 480 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481766/480-10-25-13> + Details Name: us-gaap_FairValueAdjustmentOfWarrants Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionAmount of gain (loss) on sale or disposal of property, plant and equipment assets, including oil and gas property and timber property. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Subparagraph (b) -SubTopic 10 -Topic 230 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-28> + Details Name: us-gaap_GainLossOnSaleOfPropertyPlantEquipment Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionThe increase (decrease) during the reporting period in the aggregate amount of liabilities incurred (and for which invoices have typically been received) and payable to vendors for goods and services received that are used in an entity's business. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Subparagraph (a) -SubTopic 10 -Topic 230 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-28> + Details Name: us-gaap_IncreaseDecreaseInAccountsPayable Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionThe increase (decrease) during the reporting period in amount due within one year (or one business cycle) from customers for the credit sale of goods and services. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Subparagraph (a) -SubTopic 10 -Topic 230 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-28> + Details Name: us-gaap_IncreaseDecreaseInAccountsReceivable Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionThe increase (decrease) during the reporting period in the aggregate amount of expenses incurred but not yet paid. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Subparagraph (a) -SubTopic 10 -Topic 230 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-28> + Details Name: us-gaap_IncreaseDecreaseInAccruedLiabilities Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionThe increase (decrease) during the reporting period in the assets (liabilities) created through trading commodity-based derivative instruments. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Subparagraph (a) -SubTopic 10 -Topic 230 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-28> + Details Name: us-gaap_IncreaseDecreaseInCommodityContractAssetsAndLiabilities Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionThe increase (decrease) during the reporting period in the aggregate amount due to the entity in the form of unpaid interest and dividends. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Subparagraph (a) -SubTopic 10 -Topic 230 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-28> + Details Name: us-gaap_IncreaseDecreaseInInterestAndDividendsReceivable Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - ReferencesNo definition available. + Details Name: us-gaap_IncreaseDecreaseInOperatingCapitalAbstract Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionAmount of increase (decrease) in noncurrent assets classified as other. + ReferencesReference 1: <http://www.xbrli.org/2009/role/commonPracticeRef> -Topic 230 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Subparagraph (a) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-28> + Details Name: us-gaap_IncreaseDecreaseInOtherNoncurrentAssets Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionThe increase (decrease) during the reporting period in the amount of outstanding money paid in advance for goods or services that bring economic benefits for future periods. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Subparagraph (a) -SubTopic 10 -Topic 230 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-28> + Details Name: us-gaap_IncreaseDecreaseInPrepaidExpense Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionAmount of realized and unrealized gain (loss) on investment in marketable security. + ReferencesNo definition available. + Details Name: us-gaap_MarketableSecuritiesGainLossExcludingOtherThanTemporaryImpairments Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionAmount of cash inflow (outflow) from financing activities, including discontinued operations. Financing activity cash flows include obtaining resources from owners and providing them with a return on, and a return of, their investment; borrowing money and repaying amounts borrowed, or settling the obligation; and obtaining and paying for other resources obtained from creditors on long-term credit. + ReferencesReference 1: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 230 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 24 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-24> + Details Name: us-gaap_NetCashProvidedByUsedInFinancingActivities Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - ReferencesNo definition available. + Details Name: us-gaap_NetCashProvidedByUsedInFinancingActivitiesAbstract Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionAmount of cash inflow (outflow) from investing activities, including discontinued operations. Investing activity cash flows include making and collecting loans and acquiring and disposing of debt or equity instruments and property, plant, and equipment and other productive assets. + ReferencesReference 1: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 230 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 24 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-24> + Details Name: us-gaap_NetCashProvidedByUsedInInvestingActivities Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - ReferencesNo definition available. + Details Name: us-gaap_NetCashProvidedByUsedInInvestingActivitiesAbstract Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionAmount of cash inflow (outflow) from operating activities, including discontinued operations. Operating activity cash flows include transactions, adjustments, and changes in value not defined as investing or financing activities. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 230 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-28>Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 230 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 24 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-24>Reference 3: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 230 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 25 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-25> + Details Name: us-gaap_NetCashProvidedByUsedInOperatingActivities Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: na Period Type: duration X - ReferencesNo definition available. + Details Name: us-gaap_NetCashProvidedByUsedInOperatingActivitiesAbstract Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionAmount of cash outflow from operating lease, excluding payments to bring another asset to condition and location necessary for its intended use. + ReferencesReference 1: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 842 -SubTopic 20 -Name Accounting Standards Codification -Section 45 -Paragraph 5 -Subparagraph (c) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479041/842-20-45-5>Reference 2: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 842 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph (g)(1) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478964/842-20-50-4> + Details Name: us-gaap_OperatingLeasePayments Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionThe cash outflow for loan and debt issuance costs. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Topic 230 -SubTopic 10 -Section 45 -Paragraph 15 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-15> + Details Name: us-gaap_PaymentsOffinancingCosts Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionThe cash outflow for acquisition of machinery and equipment. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 230 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 13 -Subparagraph (c) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-13> + Details Name: us-gaap_PaymentsToAcquireMachineryAndEquipment Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionAmount of cash outflow for purchase of marketable security. + ReferencesReference 1: <http://www.xbrli.org/2009/role/commonPracticeRef> -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 11 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481830/320-10-45-11> + Details Name: us-gaap_PaymentsToAcquireMarketableSecurities Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionThe cash inflow associated with the aggregate amount received by the entity through sale or maturity of marketable securities (held-to-maturity or available-for-sale) during the period. + ReferencesNo definition available. + Details Name: us-gaap_ProceedsFromSaleAndMaturityOfMarketableSecurities Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionThe cash inflow from a borrowing having initial term of repayment within one year or the normal operating cycle, if longer. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Topic 230 -SubTopic 10 -Section 45 -Paragraph 14 -Subparagraph (b) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-14> + Details Name: us-gaap_ProceedsFromShortTermDebt Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionAmount of cash inflow from exercise of option under share-based payment arrangement. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 45 -Paragraph 14 -Subparagraph (a) -SubTopic 10 -Topic 230 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482740/230-10-45-14>Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 50 -Paragraph 2A -Subparagraph (a) -SubTopic 10 -Topic 718 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2A> + Details Name: us-gaap_ProceedsFromStockOptionsExercised Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionThe consolidated profit or loss for the period, net of income taxes, including the portion attributable to the noncontrolling interest. + ReferencesReference 1: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 6 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147483443/250-10-50-6>Reference 2: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 9 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147483443/250-10-50-9>Reference 3: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 805 -SubTopic 60 -Name Accounting Standards Codification -Section 65 -Paragraph 1 -Subparagraph (g) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147476176/805-60-65-1>Reference 4: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 740 -SubTopic 323 -Name Accounting Standards Codification -Section 65 -Paragraph 2 -Subparagraph (g)(3) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478666/740-323-65-2>Reference 5: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 235 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.4-08(g)(1)(iii)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480678/235-10-S99-1>Reference 6: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 323 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph (c) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481687/323-10-50-3>Reference 7: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 825 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 28 -Subparagraph (f) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482907/825-10-50-28>Reference 8: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (b)(2) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147483443/250-10-50-1>Reference 9: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 815 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 1 -Subparagraph (f) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480175/815-40-65-1>Reference 10: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 8 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147483443/250-10-50-8>Reference 11: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 946 -SubTopic 830 -Name Accounting Standards Codification -Section 55 -Paragraph 11 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479168/946-830-55-11>Reference 12: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 946 -SubTopic 205 -Name Accounting Standards Codification -Section 45 -Paragraph 3 -Subparagraph (a) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478009/946-205-45-3>Reference 13: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section 45 -Paragraph 7 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479105/946-220-45-7>Reference 14: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 944 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-04(16)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147477520/944-220-S99-1>Reference 15: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.6-07(9)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479134/946-220-S99-1>Reference 16: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-09(1)(d)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479134/946-220-S99-3>Reference 17: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 810 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 19 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481231/810-10-45-19>Reference 18: <http://www.xbrli.org/2009/role/commonPracticeRef> -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 6 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482765/220-10-50-6>Reference 19: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(4)(i)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1A>Reference 20: <http://www.xbrli.org/2009/role/commonPracticeRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(4)(ii)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1A>Reference 21: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(4)(iii)(A)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1A>Reference 22: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(4)(iv)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1A>Reference 23: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph (SX 210.13-01(a)(5)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1A>Reference 24: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1B -Subparagraph (SX 210.13-02(a)(4)(i)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1B>Reference 25: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1B -Subparagraph (SX 210.13-02(a)(4)(iii)(A)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480097/470-10-S99-1B>Reference 26: <http://www.xbrli.org/2003/role/disclosureRef> -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1B -Subparagraph (SX 210.13-02(a)(4)(iii)(B)) -Publisher

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Codification -Section 50 -Paragraph 7 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483499/205-20-50-7Reference 31:](#) [http://www.xbrl.org/2003/role/exampleRef -Topic 810 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 4J -Publisher FASB -URI https://asc.fasb.org/1943274/2147481175/810-10-55-4JReference 32:](#) [http://www.xbrl.org/2003/role/exampleRef -Topic 810 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 4K -Publisher FASB -URI https://asc.fasb.org/1943274/2147481175/810-10-55-4KReference 33:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 1A -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482790/220-10-45-1AReference 34:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 1B -Subparagraph \(a\) -Publisher FASB -URI 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ReferencesReference 1:](#) [http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Subparagraph \(a\) -SubTopic 10 -Topic 230 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482740/230-10-45-28 + Details Name: us-gaap_ShareBasedCompensation Namespace Prefix: us-gaap_Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration XML 1.7.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Pay vs Performance Disclosure - USD \(\\$\) in Thousands 3 Months Ended 6 Months Ended Jun. 30, 2024 Mar. 31, 2024 Jan. 31, 2024 Mar. 31, 2023 Mar. 31, 2023 Jun. 30, 2023 Jun. 30, 2023 Pay vs Performance Disclosure Å Å Å Net Income \(Loss\) \\$ \(462,492\) \\$ \(14,792\) \\$ \(20,640\) \\$ \(21,726\) \\$ \(477,284\) \\$ \(42,366\) X - ReferencesReference 1:](#) [http://www.xbrl.org/2003/role/presentationRef -Publisher SEC -Name Regulation S-K -Number 229 -Section 402 -Subsection v -Paragraph 1 + Details Name: ecd_PvTable Namespace Prefix: ecd_Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionThe portion of profit or loss for the period, net of income taxes, which is attributable to the parent. + ReferencesReference 1:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 6 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-6Reference 2:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 9 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-9Reference 3:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 805 -SubTopic 60 -Name Accounting Standards Codification -Section 65 -Paragraph 1 -Subparagraph \(g\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147476176/805-60-65-1Reference 4:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 740 -SubTopic 323 -Name Accounting Standards Codification -Section 65 -Paragraph 2 -Subparagraph \(g\)\(3\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478666/740-323-65-2Reference 5:](#) [http://fasb.org/us-gaap/role/ref/legacyRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 2 -Subparagraph \(SX 210.5-03\(20\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483621/220-10-S99-2Reference 6:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 235 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.4-08\(g\)\(1\)\(iii\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480678/235-10-S99-1Reference 7:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 323 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph \(c\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481687/323-10-50-3Reference 8:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 825 -SubTopic 10 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-Paragraph 1 -Subparagraph \(f\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480175/815-40-65-1Reference 13:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 8 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-8Reference 14:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 11 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-11Reference 15:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 11 -Subparagraph \(b\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-11Reference 16:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 250 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483443/250-10-50-4Reference 17:](#) [http://www.xbrl.org/2003/role/exampleRef -Topic 946 -SubTopic 830 -Name Accounting Standards Codification -Section 55 -Paragraph 10 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479168/946-830-55-10Reference 18:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section 45 -Paragraph 7 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479105/946-220-45-7Reference 19:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.7-04\(18\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477250/944-220-S99-1Reference 20:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.6-07\(9\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-1Reference 21:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph \(SX 210.6-09\(1\)\(d\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-3Reference 22:](#) [http://www.xbrl.org/2003/role/disclosureRef -Topic 470 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1A -Subparagraph \(SX 210.13-01\(a\)\(4\)\(ii\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2](#)

only owned subsidiary of the Company. X - Definition The entire disclosure for the nature of an entity's business, major products or services, principal markets including location, and the relative importance of its operations in each business and the basis for the determination, including but not limited to, assets, revenues, or earnings. For an entity that has not commenced principal operations, disclosures about the risks and uncertainties related to the activities in which the entity is currently engaged and an understanding of what those activities are being directed toward. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef-Name Accounting Standards Codification -Topic 275 -Publisher FASB -URI https://asc.fasb.org/275/tableOfContentReference> 2:

deliverables. The collaborative arrangements do not include a right of return for any deliverable. As part of the accounting for these arrangements, the Company must develop assumptions that require judgment to determine the stand-alone selling price for each performance obligation identified in the contract. In developing the stand-alone selling price for a performance obligation, the Company considers competitor pricing for a similar or identical product, market awareness of and perception of the product, expected product life and current market trends. In general, the consideration allocated to each performance obligation is recognized when the respective obligation is satisfied either by delivering a good or providing a service, limited to the consideration that is not constrained. Non-refundable payments received before all of the relevant criteria for revenue recognition are satisfied are recorded as contract liabilities. Licenses of intellectual property If a license to the Company's intellectual property is determined to be distinct from the other promises or performance obligations identified in the contract, the Company recognizes revenue from the portion of the transaction price allocated to the license at a point in time, when the license is transferred to the customer and the customer is able to benefit from the license. Research and development services The portion of a transaction price allocated to research and development services performance obligations is deferred and recognized as revenue over time as delivery or performance of such services occurs based on the use of an input method. Customer options A customer's right to choose, at its discretion, to make a payment for additional goods or services is generally considered an option. If the Company is not presently obligated to provide and does not have a right to consideration for delivering additional goods or services, the item is considered an option. The Company evaluates the customer options for material rights, such as the ability to acquire additional goods or services for free or a discount. Optional future services that reflect their standalone selling prices do not provide the customer with a material right and, therefore, are not considered performance obligations and are accounted for as separate contracts. The optional future services do not include a material right to be accounted for as performance obligations. Milestones payments The Company's collaboration agreements include development and regulatory milestones. The Company evaluates whether the milestones are considered probable of being reached and estimates the amount to be included in the transaction price using the most likely amount method. The Company evaluates factors such as the scientific, clinical, regulatory, commercial, and other risks that must be overcome to achieve the particular milestone in making this assessment. If it is probable that a significant revenue reversal would not occur, the associated milestone value is included in the transaction price. Milestone payments that are not within the Company's control or the licensee's control, such as regulatory approvals, are not considered probable of being achieved until those approvals are received. At the end of each reporting period, the Company re-evaluates the probability of achievement of such milestones and any related constraint, and if necessary, adjusts the estimate of the overall transaction price. Any such adjustments are recorded on a cumulative catch-up basis, which would affect revenue and net loss in the period of adjustment. Royalties For sales-based royalties, including milestone payments based on the level of sales, the Company determines whether the sole or predominant item to which the royalties relate is a license. When the license is the sole or predominant item to which the sales-based royalty relates, the Company recognizes revenue at the later of: (i) when the related sales occur, or (ii) when the performance obligation to which some or all of the royalty has been allocated has been satisfied (or partially satisfied). To date, the Company has not recognized any sales-based royalty revenue resulting from the Company's collaboration agreements. The Company receives payments from its customers based on billing terms established in the contract. Up-front payments and fees are recorded as contract liabilities (e.g., deferred revenue) upon receipt or when due until the Company performs its obligations under the arrangement. In the event of an early termination of a collaboration agreement, any contract liabilities would be recognized in the period in which all our obligations under the agreement have been fulfilled. For a complete discussion of accounting for revenue, see Note 5, "Collaboration and License Agreements." Costs to Fulfill a Contract with a Customer The Company has not capitalized any costs for the three and six months ended June 30, 2024 and 2023. The Company is required to capitalize costs incurred to fulfill customer contracts. These costs are required to be amortized to expense on a systemic basis that is consistent with the transfer to the customer of the goods or services to which the asset relates, compared to previously being expensed as incurred. Elements of the costs primarily include (i) payroll and other related costs of personnel related directly to the contract activities; (ii) costs related to pre-clinical testing and clinical trials such as payments to contract research organizations ("CROs") and contract manufacturing organizations ("CMOs"), investigators, and clinical trial sites that conduct the contract activities; (iv) costs to develop the product candidates, including raw materials and supplies, product testing, depreciation, and facility-related expenses; and (v) other research and development costs. Leases The Company determines if an arrangement contains a lease at inception. For arrangements where the Company is the lessee, operating leases are included in operating lease right-of-use, or ROU assets; current operating lease liabilities; and non-current operating lease liabilities on its balance sheets. The Company currently does not have any finance leases. Operating lease ROU assets and operating lease liabilities are recognized based on the present value of the future minimum lease payments over the lease term at the commencement date. ROU assets also include any initial direct costs incurred and any lease payments made on or before the lease commencement date, less lease incentives received. The Company uses its incremental borrowing rate based on the information available at the commencement date in determining the lease liabilities as the Company's leases generally do not provide an implicit rate. The incremental borrowing rate is reevaluated upon a lease modification. The operating lease ROU asset also includes any initial direct costs and prepaid lease payments made less any lease incentives. The Company considered information available at the adoption date of Topic 842 to determine the incremental borrowing rate for leases in existence as of this date. The incremental borrowing rate used was the weighted average rate between secured and unsecured lending arrangements. Lease terms may include options to extend or terminate the lease when the Company is reasonably certain that the option will be exercised. Variable payments included in the lease agreement are expensed as incurred. Lease expense is recognized on a straight-line basis over the lease term. The Company elected to apply each of the practical expedients described in ASC Topic 842-10-65-1(f) which allow companies not to reassess: (i) whether any expired or existing agreements contain leases, (ii) the classification of any expired or existing leases, and (iii) the capitalization of initial direct costs for any existing leases. The Company also elected to apply the short-term lease measurement and recognition exemption in which ROU assets and lease liabilities are not recognized for short-term operating leases. A short-term operating lease is a lease that, at the commencement date, has a lease term of 12 months or less and does not include an option to purchase the underlying asset that the lessee is reasonably certain to exercise. The Company also elected not to separate non-lease components from lease components and instead to account for each separate lease component and the non-lease components associated with that lease component as a single lease component. Segment Information The Company manages its operations as a single segment for the purposes of assessing performance and making operating decisions. The Company's operations are focused on oncology development activities. Research and Development Costs Costs incurred in connection with research and development activities are expensed as incurred. These costs include fees paid to consultants, vendors and various entities that perform certain research and testing on behalf of the Company. The Company has acquired rights to develop and commercialize product candidates. Upfront payments that relate to the acquisition of a new product compound, as well as pre-commercial milestone payments, are immediately expensed as acquired in-process research and development in the period in which they are incurred, provided that the new product compound did not also include processes or activities that would constitute a "business" as defined under U.S. GAAP, and the product candidate has not achieved regulatory approval for marketing and, absent obtaining such approval, has no established alternative future use. Milestone payments made to third parties subsequent to regulatory approval which meet the capitalization criteria would be capitalized as intangible assets and amortized over the estimated remaining useful life of the related product. If the conditions enabling capitalization of development costs as an asset have not yet been met, all development expenditures are recognized in profit or loss when incurred. Stock-based Compensation The Company recognizes compensation cost for grants of employee stock options using a fair-value measurement method, that is recognized in operating results as compensation expense based on the fair value over the requisite service period of the awards. Forfeitures are recorded as they occur instead of estimating forfeitures that are expected to occur. The Company determines the fair value of stock-based awards that are based only on a service condition using the Black-Scholes option-pricing model which uses both historical and current market data to estimate fair value. The method incorporates various assumptions such as the risk-free interest rate, volatility, dividend yield, and expected life of the options. The Company determines the fair value of stock-based awards that are based on both a service condition and achievement of the first to occur of a market or performance condition using a Monte Carlo simulation. Income Taxes The Company accounts for income taxes using the asset and liability method, which requires the recognition of deferred tax assets and liabilities for the expected future tax consequences of events that have been recognized in the consolidated financial statements or in the Company's tax returns. The difference between the financial statement and tax basis of assets and liabilities is determined annually. Deferred income tax assets and liabilities are computed for those differences that have future tax consequences using the currently enacted tax laws and rates that apply to the years in which they are expected to affect taxable income. The Company assesses the likelihood that its deferred tax assets will be recovered from future taxable income and, to the extent it believes, based upon the weight of available evidence, that it is more likely than not that all or a portion of the deferred tax assets will not be realized, a valuation allowance is established through a charge to income tax expense. The Company uses a recognition threshold and measurement attribute for the financial statement recognition and measurement of a tax position taken, or expected to be taken, in a tax return. The provision for income taxes includes the effects of any resulting tax reserves, or unrecognized tax benefits, that are considered appropriate. The Company's policy is to record interest and penalties related to income taxes as part of the tax provision. Returns for tax years beginning with those filed for the period ended December 31, 2018 are open to federal and state tax examination. Recent Accounting Pronouncements In November 2023, the FASB issued Topic 280 "Improvements to Reportable Segment Disclosures" which expands annual and interim disclosure requirements for reportable segments, primarily through enhanced disclosures about significant segment expenses. This guidance is effective for our annual periods beginning January 1, 2024, and for interim periods beginning January 1, 2025, with early adoption permitted. We are currently evaluating the potential effect that the updated standard will have on our financial statement disclosures. In December 2023, the FASB issued Topics 740 "Improvements to Income Tax Disclosures" to expand the disclosure requirements for income taxes, specifically related to the rate reconciliation and income taxes paid. This guidance is effective for our annual periods beginning January 1, 2025, with early adoption permitted. We are currently evaluating the potential effect that the updated standard will have on our financial statement disclosures. X - References No definition available. + Details Name: us-gaap_AccountingPoliciesAbstract Namespace Prefix: us-gaap_Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - Definition The entire disclosure for all significant accounting policies of the reporting entity. + References Reference 1: <http://www.xbrl.org/2003/role/disclosureRef> - Topic 235 - Sub-Topic 10 - Name Accounting Standards Codification - Section 50 - Paragraph 1 - Publisher FASB - URI <https://asc.fasb.org/1943274/2147483426/235-10-50-1> Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> - Topic 235 - Name Accounting Standards Codification - Publisher FASB - URI <https://asc.fasb.org/235/tableOfContent> + Details Name: us-gaap_SignificantAccountingPoliciesTextBlock Namespace Prefix: us-gaap_Data Type: dtr-types:textBlock Item Type Balance Type: na Period Type: duration XML 18 R11.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Acquisition 6 Months Ended Jun. 30, 2024 Business Combinations [Abstract] A Acquisition 3. Acquisition On the Acquisition Date, the Company completed the Acquisition of AnHeart. Pursuant to the terms of the AnHeart Merger Agreement, at the effective time of the First Merger (as defined in the Merger Agreement) (the "First Effective Time"), the Company issued to AnHeart securityholders (i) approximately 27,646,255 shares of Class A Common Stock, (ii) 851,202 shares of Series A Non-Voting Convertible Preferred Stock of the Company, and (iii) the Consideration Warrants. The shares of our Series A Non-Voting Convertible Preferred Stock are automatically convertible into an aggregate of approximately 85,120,200 shares of Class A Common Stock upon the approval of such conversion by the Company's stockholders in accordance with the rules of the New York Stock Exchange. Since the preferred stock is not mandatorily redeemable and the exception for a deemed liquidation event was not met, all of the outstanding convertible preferred stock is classified as mezzanine equity in the balance sheet. The Consideration Warrants are restricted with respect to the exercise and transfer thereof until receipt of such stockholder approval and otherwise have terms identical to those of the Company's outstanding Public Warrants. At the First Effective Time, (i) each option to purchase shares of Ordinary Shares (as defined in the Merger Agreement) of AnHeart (an "AnHeart Option") held by a Continuing Company Service Provider (as defined in the Merger Agreement), whether or not vested, was assumed and converted into an option to purchase Class A Common Stock (each such option, an "Assumed Option"), and (ii) each restricted stock unit reflecting the right to receive Ordinary Shares of AnHeart (an "AnHeart RSU") held by a Continuing Company Service Provider (as defined in the Merger Agreement) was assumed and became a restricted stock unit with respect to a number of shares of Class A Common Stock (each such restricted stock unit, an "Assumed RSU"). Each Assumed Option and Assumed RSU is subject to the same terms and conditions (including vesting and exercise schedule) as were applicable to the corresponding AnHeart Option or AnHeart RSU immediately prior to the First Effective Time, subject to limited exceptions set forth in the Merger Agreement. Any other AnHeart Option and AnHeart RSU that remained unexercised and outstanding as of immediately prior to the First Effective Time was canceled without payment. From and after the Effective Time: (a) the number of shares of Class A Common Stock subject to each Assumed Option was determined by multiplying (A) the number of Ordinary Shares of AnHeart that are subject to such AnHeart Option immediately prior to the First Effective Time, by (B) the Equity Award Exchange Ratio (as defined in the Merger Agreement) and rounding down to the nearest whole number of shares; (b) the per-share exercise price for Class A Common Stock issuable upon exercise of each Assumed Option was determined by dividing (A) the per-share exercise price of each AnHeart Option, as in effect immediately prior to the First Effective Time, by (B) the Equity Award Exchange Ratio and rounding up to the nearest whole cent; and (c) the number of shares of Class A Common Stock subject to each Assumed RSU was determined by multiplying (A) the number of Ordinary Shares of AnHeart that are subject to such AnHeart RSU immediately prior to the First Effective Time, by (B) 1.7551 and rounding down to the nearest whole number of shares. The final allocable purchase price consists of the following (in thousands): Class A Common Stock (1) \$ 89,297 A Series A Non-Voting Convertible Preferred Stock (2) A \$ 274,938 A Consideration Warrants (3) A \$ 764 A Assumed share-based awards (4) A \$ 24,818 A Acquisition costs A \$ 7,434 A Total allocable purchase price A \$ 397,251 A A \$ (1) 27,646,255 shares issued at \$3.23 per share as of the Acquisition Date. (2) 851,202 shares issued at as-converted value of \$323 per share. (3) 2,893,731 warrants issued at the estimated fair value of \$0.2641 per warrant. (4) The portion recorded in consideration transferred represents the fair value of replacement awards (Assumed Options & Assumed RSUs) attributable to precombination services. We accounted for the transaction as an asset acquisition since the lead asset, talrectinib, represented substantially all of the fair value of the gross assets acquired, which exclude cash acquired. At the Acquisition Date, we recorded a \$425.1 million charge representing an acquired in-process research and development asset with no alternative future use in acquired in-process research and development expenses, of which the \$425.1 million net cash consideration is presented as a cash outflow from operating activities. In connection with this acquisition, we recorded \$450.9 million of assets acquired primarily consisting of cash and investments. Liabilities assumed were approximately \$53.7 million. X - Definition The entire disclosure for a business combination (or series of individually immaterial business combinations) completed during the period, including background, timing, and recognized assets and liabilities. The disclosure may include leverage buyout transactions (as applicable). + References Reference 1: <http://www.xbrl.org/2003/role/disclosureRef> - Topic 805 - Sub-Topic 20 - Name Accounting Standards Codification - Section 50 - Paragraph 5 - Subparagraph (b) - Publisher FASB - URI <https://asc.fasb.org/1943274/2147479907/805-20-50-5> Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> - Topic 805 - Name Accounting Standards Codification - Publisher FASB - URI <https://asc.fasb.org/805/tableOfContent> + Details Name: us-gaap_BusinessCombinationDisclosureTextBlock Namespace Prefix: us-gaap_Data Type: dtr-types:textBlock Item Type Balance Type: na Period Type: duration X - References No definition available. + Details Name: us-gaap_BusinessCombinationsAbstract Namespace Prefix: us-gaap_Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration XML 19 R12.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Fair Value Measurements and Marketable Securities Available-for-Sale 6 Months Ended Jun. 30, 2024 Fair Value Disclosures [Abstract] A Fair Value Measurements and Marketable Securities Available for Sale 4. Fair Value Measurements and Marketable Securities Available-for-Sale The Company provides disclosure of financial assets and financial liabilities that are carried at fair value based on the price that would be received upon sale of an asset or paid to transfer a liability in an orderly transaction between market participants at the measurement date. Fair value measurements may be classified based on the amount of subjectivity associated with the inputs to fair valuation of these assets and liabilities using the following three levels: Level 1 Inputs are unadjusted quoted prices in active markets for identical assets or liabilities that the Company has the ability to access at the measurement date. Level 2 Inputs include quoted prices for similar assets and liabilities in active markets, quoted prices for identical or similar assets or liabilities in markets that are not active, inputs other than quoted prices that are observable for the asset or liability (i.e., interest rates, yield curves, etc.) and inputs that are derived principally from or corroborated by observable market data by correlation or other means (market corroborated inputs). Level 3 Inputs are unobservable inputs that reflect the Company's estimates of the assumptions that market participants would use in pricing the asset or liability. The Company develops these inputs based on the best information available, including its own data. The following table presents information about the Company's marketable securities as of June 30, 2024 and December 31, 2023 and the Warrant liability as of June 30, 2024 and December 31, 2023,

measured at fair value on a recurring basis, and indicates the fair value hierarchy of the valuation techniques the Company utilized to determine such fair value. The Company's Warrant liabilities are included within the Level 1 and Level 3 fair value hierarchy. The fair value of the Consideration, Public and Forward Purchase Warrants is determined using the closing price of the warrants on the NYSE market. The fair value of the Private Placement Warrants is determined using the Black-Scholes option pricing formula. The primary unobservable input utilized in determining the fair value of the Private Warrants is the expected volatility. The expected volatility was estimated considering observable Public Warrant pricing, the Company's own historical volatility and the volatility of guideline public companies. There have not been any transfers between the levels during the periods.

	June 30, 2024	December 31, 2023
Financial assets:	\$ 7,492	\$ 7,492
Cash equivalents:	\$ 19,556	\$ 19,556
Corporate bonds	\$ 2,138	\$ 2,138
Certificate of deposits	\$ 21,694	\$ 21,694
Commercial paper	\$ 36,555	\$ 36,555
U.S. government and government agency securities	\$ 408,996	\$ 408,996
Corporate bonds	\$ 75,639	\$ 75,639
Financial liabilities:	\$ 1,441	\$ 1,363
Warrants	\$ 78	\$ 78
December 31, 2023	\$ 20,574	\$ 20,574
Financial assets:	\$ 7,492	\$ 7,492
Cash equivalents:	\$ 19,556	\$ 19,556
Corporate bonds	\$ 2,138	\$ 2,138
Certificate of deposits	\$ 21,694	\$ 21,694
Commercial paper	\$ 36,555	\$ 36,555
U.S. government and government agency securities	\$ 408,996	\$ 408,996
Corporate bonds	\$ 75,639	\$ 75,639
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Financial liabilities:	\$ 1,441	\$ 1,363
Warrants	\$ 78	\$ 78
December 31, 2023	\$ 2	

contracts for the three months ended June 30, 2024. The costs incurred to fulfill customer contracts were capitalized and amortized to cost of revenue on a systemic basis that is consistent with the transfer of R&D services to the customer to which the asset relates. For the three months ended June 30, 2024, \$1.3 million of costs incurred to fulfill customer contracts were capitalized and expensed in the same period. There were no balances related to the costs incurred to fulfill customer contracts as of June 30, 2024. X - DefinitionCollaboration and Licence Agreement [Abstract] + ReferencesNo definition available. + Details Name: nuvb CollaborationAndLicenceAgreementAbstract Namespace Prefix: nuvb Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionCollaborationAnd Licence Agreements TextBlock + ReferencesNo definition available. + Details Name: nuvb CollaborationAndLicenceAgreementsTextblock Namespace Prefix: nuvb Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration XML 21 R14.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Borrowings 6 Months Ended Jun. 30, 2024 Subordinated Borrowings [Abstract] A Borrowings BorrowingIn 2020, AnHeart entered into loan agreements with Bank of Hangzhou to obtain short-term borrowings to supplement its working capital. As of June 30, 2024, the outstanding balance net of repayments was \$7.0 million. For the three months ended June 30, 2024, the Company had drawn down \$7.0 million and repaid \$5.6 million. The fixed interest rate of these borrowings was 4.05% per annum. In 2022, AnHeart entered into loan agreements with China Merchants Bank to obtain short-term borrowings to supplement its working capital. As of June 30, 2024, the outstanding balance was \$2.8 million. For the three months ended June 30, 2024, there were no borrowings or repayments. The fixed interest rate of these borrowings was 3.90% per annum. On April 4, 2023, AnHeart entered into a Loan and Security Agreement ("2023 SSVB Agreement") with Shanghai Pudong Development Bank in Silicon Valley for up to 40.0 million RMB or equivalent in optional currency USD term loans ("SSVB Loan"). The SSVB Loan consists of a short-term working capital loan of 20 million RMB and a long-term loan of 20 million RMB, which matures on April 4, 2024 and April 4, 2025 respectively, at which time all outstanding balances are due. Draws on the line of credit for the short-term loans are payable on the maturity date of the SSVB Loan. Draws on the line of credit for the long-term loans are payable in equal instalments monthly over the shorter of 24 months or from the date the draw took place till the SSVB loan matures. Outstanding balances will bear interest at a fixed rate of 4.25%. As of June 30, 2024, the outstanding line of credit balance net of repayments was \$1.8 million. For the three months ended June 30, 2024, the Company had repaid \$ 0.7 million. X - ReferencesNo definition available. + Details Name: us-gaap SubordinatedBorrowingsAbstract Namespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionThe entire disclosure for borrowings under subordinated debt agreements that qualify as available in computing net capital under SEC's uniform net capital rule, including restrictive covenants, collateral, interest rates and due dates, amounts due by date and amount owed in total. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 942 -SubTopic 470 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147478878/942-470-45-1> + Details Name: us-gaap SubordinatedBorrowingsDisclosureTextBlock Namespace Prefix: us-gaap Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration XML 22 R15.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Leases 6 Months Ended Jun. 30, 2024 Leases [Abstract] A LeasesOur principal executive office is located in New York, New York, where we lease approximately 7,900 square feet of office space under a lease that terminates in 2027, with an option for the Company to extend the lease for an additional five years which is not reasonably assured of exercise. We also occupy approximately 25,139 square feet of office space in San Francisco, California, under a lease that terminates in 2025 and a total of approximately 1,582 square meters of office space in the Peopleleâ€™s Republic of China, in the cities of Beijing, Guangzhou, Hangzhou, Shanghai and Yantai, under leases that terminate in 2024 through 2026.Operating lease expense was \$0.5 million and \$0.4 million for the three months ended June 30, 2024 and 2023, respectively, and \$1.0 million and \$0.7 million for the six months ended June 30, 2024 and 2023, respectively. Expense related to variable leases were not significant for the three and six months ended June 30, 2024 and 2023. Operating cash flows for the six months ended June 30, 2024 and 2023 included \$1.1 million and \$0.7 million for operating leases, respectively. The following table presents the future minimum lease analysis of the Company's operating lease liabilities showing the aggregate lease payments as of June 30, 2024.A A A June 30, 2024 A A A (In thousands) A 2024 (remaining 6 months) A \$ 1,095 A 2025 A 1,416 A 2026 A 629 A 2027 and thereafter A 81 A Total undiscounted lease payments A 3,221 A Less: imputed interest A (173) A Total operating lease liabilities A \$ 3,048 A The weighted average incremental borrowing rate used to determine the operating lease liabilities was 6.81%. The Company's weighted average remaining lease term was 1.81 years as of June 30, 2024. X - ReferencesNo definition available. + Details Name: us-gaap LeasesAbstract Namespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionThe entire disclosure for operating leases of lessee. Includes, but is not limited to, description of operating lease and maturity analysis of operating lease liability. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 842 -SubTopic 20 -Name Accounting Standards Codification -Publisher FASB -URI <https://asc.fasb.org/842-20/tableOfContent> + Details Name: us-gaap LesseeOperatingLeasesTextBlock Namespace Prefix: us-gaap Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration XML 23 R16.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Net Loss Per Share 6 Months Ended Jun 30, 2024 Earnings Per Share [Abstract] A Net Loss per Share 8. Net Loss per Share Net loss per common share is computed using the two-class method required due to the participating nature of the convertible preferred stock. Although the convertible preferred stock are participating securities, such securities do not participate in net losses and therefore do not impact the Companyâ€™s net loss from continuing operations per share calculation as of June 30, 2024.Basic loss per share is computed by dividing net loss by the weighted-average number of shares of Class A and Class B Common Stock outstanding. Diluted loss per share reflects the potential dilution that could occur if the stock options to issue Common Stock were exercised. The Company had a net loss in all periods presented thus the dilutive net loss per common share is the same as the basic net loss per common share as the effect of any options or conversions is anti-dilutive.The earnings per share amounts are the same for the different classes of Common Stock because the holders of each class are legally entitled to equal per share distributions whether through dividends or liquidation.The following securities outstanding at June 30, 2024 and 2023 have been excluded from the calculation of weighted average shares outstanding:A A A As of June 30, A A A 2024 A 2023 A Warrants A 8,681,193 A A 5,787,462 A Class A Common Stock options A 52,692,948 A A 33,208,484 A Restricted stock units A 2,161,352 A A A â€” A Convertible preferred stock A 85,120,200 A A A â€” A X - ReferencesNo definition available. + Details Name: us-gaap EarningsPerShareAbstract Namespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionThe entire disclosure for earnings per share. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (c) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482662/260-10-50-1-Reference 2>: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 260 -Name Accounting Standards Codification -Publisher FASB -URI <https://asc.fasb.org/260/tableOfContent>Reference 3: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482662/260-10-50-2-Reference 4>: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 260 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147482662/260-10-50-3> + Details Name: us-gaap EarningsPerShareTextBlock Namespace Prefix: us-gaap Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration XML 24 R17.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Accumulated Other Comprehensive Income 6 Months Ended Jun. 30, 2024 Statement of Other Comprehensive Income [Abstract] A Accumulated Other Comprehensive Income 9. Accumulated Other Comprehensive Income (Loss)The following table presents a rollforward of the changes in accumulated other comprehensive income (losses) for the six months ended June 30, 2024 and 2023, which is all attributable to unrealized gains (losses) on available-for-sale securities. All amounts are net of tax.A A A 2024 A A 2023 A Balance at beginning of period A \$ 181 A A \$ (5,526) A Unrealized gain (loss) A \$ (1,601) A A \$ 936 A Amount reclassified for realized gain on marketable securities A 6 A A A 195 A Currency translation adjustment A 148 A A A â€” A Balance at end of period A \$ (1,266) A A \$ (4,395) X - DefinitionThe entire disclosure for tabular information relating to Other Comprehensive Income (OCI) as is applicable to noncontrolling interests. This text block may also include OCI relative to the filing entity, the aforementioned noncontrolling interest OCI, as well as OCI on a consolidated basis. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 810 -Name Accounting Standards Codification -Publisher FASB -URI <https://asc.fasb.org/810/tableOfContent> + Details Name: us-gaap OtherComprehensiveIncomeNoncontrollingInterestTextBlock Namespace Prefix: us-gaap Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - ReferencesNo definition available. + Details Name: us-gaap StatementOfOtherComprehensiveIncomeAbstract Namespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration XML 25 R18.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Stock-Based Compensation 6 Months Ended Jun. 30, 2024 Share-Based Payment Arrangement [Abstract] A Stock-Based Compensation 10. Stock-Based CompensationThe 2021 Equity Incentive Plan In March 2019, the Company adopted the 2019 Equity Incentive Plan or (â€œ2019 Planâ€), which provided for the grant of options, stock appreciation rights, restricted stock, and other stock awards. In January 2021, our board of directors adopted the 2021 Equity Incentive Plan (the â€œ2021 Planâ€). The 2021 Plan was approved by our stockholders in February 2021 and became effective immediately upon the Closing Date of the Merger. Shares available for future issuance under the 2019 Plan were canceled.Awards. The 2021 Plan provides for the grant of incentive stock options (â€œISOsâ€), within the meaning of Section 422 of the Code to employees, including employees of any parent or subsidiary, and for the grant of nonstatutory stock options (â€œNSOsâ€), stock appreciation rights, restricted stock awards, restricted stock unit awards, performance awards and other forms of awards to employees, directors and consultants, including employees and consultants of our affiliates. Authorized Shares. The maximum number of shares of Class A Common Stock reserved for issuance upon adoption of the 2021 Plan was 50,684,047 shares of Class A Common Stock. The number of shares of Class A Common Stock reserved for issuance under the 2021 Plan will automatically increase on January 1 of each year, starting on January 1, 2022 through January 1, 2031, in an amount equal to (1) 4.0% of the total number of shares of Class A Common Stock and Class B Common Stock outstanding or issuable upon conversion or exercise of outstanding instruments on December 31 of the preceding year, or (2) a lesser number of shares of Class A Common Stock determined by our board of directors prior to the date of the increase. The maximum number of shares of Class A Common Stock that may be issued on the exercise of ISOs under the 2021 Plan is three times the number of shares available for issuance upon the 2021 Plan becoming effective or 152,052,141 shares.The Employee Stock Purchase Plan In January 2021, our board of directors adopted the 2021 Employee Stock Purchase Plan (the â€œESPPâ€). The ESPP was approved by our stockholders in February 2021 and became effective immediately upon the Closing Date of the Merger. Share Reserve. The maximum number of shares of Class A Common Stock reserved for issuance upon the adoption of the 2021 ESPP was 4,750,354 shares of Class A Common Stock. The number of shares of Class A Common Stock reserved for issuance under the 2021 ESPP will automatically increase on January 1st of each year, beginning on January 1, 2022 and continuing through and including January 1, 2031, by 1.0% of the total number of shares of Class A Common Stock and Class B Common Stock outstanding or issuable upon conversion or exercise of outstanding instruments on December 31st of the preceding calendar year or such lesser number of shares of Class A Common Stock as determined by our board of directors. Shares subject to purchase rights granted under the 2021 ESPP that terminate without having been exercised in full will not reduce the number of shares available for issuance under the 2021 ESPP. The stock-based compensation expense included in the Companyâ€™s condensed statement of operations and comprehensive loss for the three and six months ended June 30, 2024 and 2023 is as follows (in thousands):A A A Three Months Ended June 30, A A Six Months Ended June 30, A A 2024 A A 2023 A A 2024 A 2023 A Research and development A \$ 4,190 A A \$ 2,420 A A \$ 6,482 A A \$ 4,431 A General and administrative A 5,379 A A 3,103 A A 8,128 A A 5,929 A A \$ 9,569 A A \$ 5,523 A A \$ 14,610 A A \$ 10,360 A A The following table summarizes stock option activity under the 2019 Plan, 2021 Plan and assumed options in the AnHeart Acquisition for the six months ended June 30, 2024. A A Shares A Outstanding at December 31, 2023 A A 30,649,239 A Options assumed for the Acquisition of AnHeart A 13,742,239 A Granted A A 11,322,679 A Exercised A A (1,798,190) A Forfeited or canceled A A (1,223,019) A Outstanding at June 30, 2024 A A 52,692,948 A Exercisable at June 30, 2024 A A 23,909,107 A A The weighted average exercise price of all outstanding options as of June 30, 2024 was \$2.80 per share. The following table summarizes the activity for the restricted stock units assumed in the AnHeart Acquisition for the six months ended June 30, 2024. A A Restricted Stock A A A Units A Non-vested at December 31, 2023 A A A RSUs assumed for the Acquisition of AnHeart A A 2,201,694 A Granted A A â€” A Vested A A (40,342) A Forfeited or canceled A A â€” A Non-vested at June 30, 2024 A A 2,161,352 A A A A X - DefinitionThe entire disclosure for share-based payment arrangement. + ReferencesReference 1: <http://www.xbrl.org/2003/role/exampleRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (a)(1) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2-Reference 2>: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -Name Accounting Standards Codification -Publisher FASB -URI <https://asc.fasb.org/718/tableOfContent>Reference 3: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (a) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2-Reference 4>: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (h)(2) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2-Reference 5>: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (h)(2)(i) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2-Reference 6>: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (l) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> + Details Name: us-gaap DisclosureOfCompensationRelatedCostsShareBasedPaymentsTextBlock Namespace Prefix: us-gaap Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - ReferencesNo definition available. + Details Name: us-gaap DisclosureOfCompensationRelatedCostsSharebasedPaymentsAbstract Namespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration XML 26 R19.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Warrants 6 Months Ended Jun. 30, 2024 Warrant Disclosure [Abstract] A Warrants 11. WarrantsFollowing the Merger, there were 5,787,472 warrants to purchase Common Stock outstanding, consisting of 4,791,639 Public Warrants, 162,500 Private Placement Warrants and 833,333 Forward Purchase Warrants. Each whole warrant entitles the registered holder to purchase one share of our Class A Common Stock at a price of \$11.50 per share. Pursuant to the warrant agreement, a warrant holder may exercise its warrants only for a whole number of shares of our Class A Common Stock. Following the Acquisition, the Company issued 2,893,731 Consideration Warrants. The Consideration Warrants are restricted with respect to the exercise and transfer thereof until receipt of such stockholder approval and otherwise have terms identical to those of the Companyâ€™s outstanding Public Warrants.At June 30, 2024, there were an aggregate of 8,681,193 warrants outstanding.The Company concluded that the Consideration Warrants, Public Warrants, Private Warrants and Forward Purchase Warrants do not meet the conditions to be classified in equity. The warrants were recorded at fair value with subsequent changes in fair value reflected in earnings (see Note 4). The change in fair value resulted in a gain of \$1.1 million during the three months ended June 30, 2024.The fair value of Consideration, Public and Forward Purchase Warrants is determined using the closing price of the warrants on the NYSE market and the related Warrant liability is included in Level 1 fair value measurements. The Company utilizes the Black-Scholes option pricing formula to determine the fair value of the Private Warrants at each reporting period, with changes in fair value recognized in the statement of operations. The estimated fair value of the warrant liability for the Private Warrants is determined using Level 3 inputs. Inherent in a binomial options pricing model are assumptions related to expected share-price volatility, expected life, risk-free interest rate and dividend yield. The annualized volatility of the Warrant was based on a calibration to the publicly traded warrant price as of the valuation date. The risk-free interest rate was estimated using linear interpolation assuming a term consistent with the time until the warrants expire, and yield information was based on U.S. Treasury Constant Maturities. The expected life of the warrants is assumed to be equivalent to their remaining contractual term. The dividend rate is based on the historical rate, which the Company anticipates remaining at zero.The aforementioned warrant liabilities are not subject to qualified hedge accounting.There were no transfers between Levels 1, 2 or 3 during the period ended June 30, 2024.The following table provides quantitative assumptions regarding Level 3 fair value measurements:A A June 30, 2024 A A December 31, 2023 A Stock price A \$ 2.92 A A \$ 1.51 A Strike price A \$ 11.50 A A \$ 11.50 A Term (in years) A 1.6 A A 2.1 A Volatility A 98.1 A A 82.7 % Risk-free rate A A 4.9 % A A 4.1 % Dividend yield A A 0.0 % A A 0.0 % The Company determined the following fair values for the outstanding warrants (in thousands):A A June 30,2024 A Public Warrants A \$ 767 A Private Placement Warrants A 78 A Forward Purchase Warrants A 133 A Consideration Warrants A 463 A Total A \$ 1,441 A A The following presents changes in liabilities classified in Level 3 of the fair value hierarchy for the six months ended June 30, 2024 (in thousands): A Six Months Ended A June 30, 2024 A Beginning balance A \$ 15 A Change in fair value of Private Warrants liability recognized in earnings A 63 A Ending balance A 78 A A A X - DefinitionWarrant disclosure. + ReferencesNo definition available. + Details Name: nuvb WarrantDisclosureAbstract Namespace Prefix: nuvb Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionWarrant disclosure. + ReferencesNo definition available. + Details Name: nuvb WarrantDisclosureTextBlock Namespace Prefix: nuvb Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration XML 27

R20.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Commitments and Contingencies 6 Months Ended Jun. 30, 2024 Commitments and Contingencies Disclosure [Abstract] A Commitments and Contingencies 12. Commitments and ContingenciesCommitmentsThe Company leases its office space under non-cancellable operating lease agreements. These leases also require the Company to pay real estate taxes and other operational expenses associated with the leased locations and are included in rent expense. The effect of graduating rents, net of the rent credits, is being amortized over the life of the leases so as to result in equal monthly rent expense over the lease term. The Company has standby letters of credit with banks in the aggregate amount of \$0.6 million which serve as security for the New York and San Francisco spaces operating leases. The standby letters of credit automatically renew annually.ContingenciesFrom time to time, the Company is involved in routine litigation that arises in the ordinary course of business. There are no pending significant legal proceedings to which the Company is a party, for which management believes the ultimate outcome would have a material adverse effect on the Company's financial position. X - ReferencesNo definition available. + Details Name: us-gaap CommitmentsAndContingenciesDisclosureAbstract Namespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionThe entire disclosure for commitments and contingencies. + ReferencesReference 1: <http://www.xbrl.org/2009/role/commonPracticeRef -Topic 405 -SubTopic 30 -Name Accounting Standards Codification -Publisher FASB -URI https://asc.fasb.org/405-30/tableOfContentReference> 2: [http://www.xbrl.org/2003/role/disclosureRef -Topic 440 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482648/440-10-50-4Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 440 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482648/440-10-50-4Reference) 3: <http://www.xbrl.org/2009/role/commonPracticeRef -Topic 450 -Name Accounting Standards Codification -Publisher FASB -URI https://asc.fasb.org/450/tableOfContentReference> 4: [http://www.xbrl.org/2003/role/disclosureRef -Topic 954 -SubTopic 440 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478522/954-440-50-1Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 954 -SubTopic 440 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478522/954-440-50-1Reference) 5: [http://www.xbrl.org/2003/role/disclosureRef -Topic 440 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph \(c\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482648/440-10-50-4Reference](http://www.xbrl.org/2003/role/disclosureRef -Topic 440 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482648/440-10-50-4Reference) 6:

2018, the FASB issued Topic 808, "Collaborative Arrangements" ("ASC 808"): Clarifying the Interaction between ASC 808 and ASC 606. This update clarifies that certain transactions between participants in a collaborative arrangement should be accounted for under ASC 606 when the counterparty is a customer and precludes an entity from presenting consideration from a transaction in a collaborative arrangement as revenue from contracts with customers if the counterparty is not a customer for that transaction. The Company adopted this standard for all periods presented. The Company enters into collaborative arrangements for the research and development, manufacture and/or commercialization of drug products and drug candidates. The Company assessed and determined that none of the collaboration agreements entered into during the periods presented were within the scope of ASC 808, as all of the agreements did not involve active participation by both parties in a joint research activity, and therefore did not qualify as collaborative arrangements under ASC 808. The Company has determined that all the elements of the above collaborations are reflective of a vendor-customer relationship and therefore within the scope of ASC 606. In determining the appropriate amount of revenue to be recognized as it fulfills its obligations under each of its agreements, the Company performs the five-step model under ASC 606 noted above. The Company recognizes revenue to depict the transfer of control of promised goods or services to customers in an amount that reflects the consideration to which the entity expects to receive in exchange for those goods or services. The Company's collaborative arrangements may contain more than one unit of account, or performance obligation, including grants of licenses to intellectual property rights, agreements to provide research and development services and other deliverables. The collaborative arrangements do not include a right of return for any deliverable. As part of the accounting for these arrangements, the Company must develop assumptions that require judgment to determine the stand-alone selling price for each performance obligation identified in the contract. In developing the stand-alone selling price for a performance obligation, the Company considers competitor pricing for a similar or identical product, market awareness of and perception of the product, expected product life and current market trends. In general, the consideration allocated to each performance obligation is recognized when the respective obligation is satisfied either by delivering a good or providing a service, limited to the consideration that is not constrained. Non-refundable payments received before all of the relevant criteria for revenue recognition are satisfied are recorded as contract liabilities. Licenses of intellectual property If a license to the Company's intellectual property is determined to be distinct from the other promises or performance obligations identified in the contract, the Company recognizes revenue from the portion of the transaction price allocated to the license at a point in time, when the license is transferred to the customer and the customer is able to benefit from the license. Research and development services The portion of a transaction price allocated to research and development services performance obligations is deferred and recognized as revenue over time as delivery or performance of such services occurs based on the use of an input method. Customer options A customer's right to choose, at its discretion, to make a payment for additional goods or services is generally considered an option. If the Company is not presently obligated to provide and does not have a right to consideration for delivering additional goods or services, the item is considered an option. The Company evaluates the customer options for material rights, such as the ability to acquire additional goods or services for free or a discount. Optional future services that reflect their standalone selling prices do not provide the customer with a material right and, therefore, are not considered performance obligations and are accounted for as separate contracts. The optional future services do not include a material right to be accounted for as performance obligations. Milestones payments The Company's collaboration agreements include development and regulatory milestones. The Company evaluates whether the milestones are considered probable of being reached and estimates the amount to be included in the transaction price using the most likely amount method. The Company evaluates factors such as the scientific, clinical, regulatory, commercial, and other risks that must be overcome to achieve the particular milestone in making this assessment. If it is probable that a significant revenue reversal would not occur, the associated milestone value is included in the transaction price. Milestone payments that are not within the Company's control or the licensee's control, such as regulatory approvals, are not considered probable of being achieved until those approvals are received. At the end of each reporting period, the Company re-evaluates the probability of achievement of such milestones and any related constraint, and if necessary, adjusts the estimate of the overall transaction price. Any such adjustments are recorded on a cumulative catch-up basis, which would affect revenue and net loss in the period of adjustment. Royalties For sales-based royalties, including milestone payments based on the level of sales, the Company determines whether the sole or predominant item to which the royalties relate is a license. When the license is the sole or predominant item to which the sales-based royalty relates, the Company recognizes revenue at the later of: (i) when the related sales occur, or (ii) when the performance obligation to which some or all of the royalty has been allocated has been satisfied (or partially satisfied). To date, the Company has not recognized any sales-based royalty revenue resulting from the Company's collaboration agreements. The Company receives payments from its customers based on billing terms established in the contract. Up-front payments and fees are recorded as contract liabilities (e.g., deferred revenue) upon receipt or when due until the Company performs its obligations under the arrangement. In the event of an early termination of a collaboration agreement, any contract liabilities would be recognized in the period in which all our obligations under the agreement have been fulfilled. For a complete discussion of accounting for revenue, see Note 5, "Collaboration and License Agreements." Costs to Fulfill a Contract with a Customer Costs to Fulfill a Contract with a Customer The Company has not capitalized any costs for the three and six months ended June 30, 2024 and 2023. The Company is required to capitalize costs incurred to fulfill customer contracts. These costs are required to be amortized to expense on a systematic basis that is consistent with the transfer to the customer of the goods or services to which the asset relates, compared to previously being expensed as incurred. Elements of the costs primarily include (i) payroll and other related costs of personnel related directly to the contract activities; (ii) costs related to pre-clinical testing and clinical trials such as payments to contract research organizations ("CROs") and contract manufacturing organizations ("CMOs"), investigators, and clinical trial sites that conduct the contract activities; (iv) costs to develop the product candidates, including raw materials and supplies, product testing, depreciation, and facility-related expenses; and (v) other research and development costs. Leases Leases The Company determines if an arrangement contains a lease at inception. For arrangements where the Company is the lessee, operating leases are included in operating lease right-of-use, or ROU assets; current operating lease liabilities; and non-current operating lease liabilities on its balance sheets. The Company currently does not have any finance leases. Operating lease ROU assets and operating lease liabilities are recognized based on the present value of the future minimum lease payments over the lease term at the commencement date. ROU assets also include any initial direct costs incurred and any lease payments made on or before the lease commencement date, less lease incentives received. The Company uses its incremental borrowing rate based on the information available at the commencement date in determining the lease liabilities as the Company's leases generally do not provide an implicit rate. The incremental borrowing rate is reevaluated upon a lease modification. The operating lease ROU asset also includes any initial direct costs and prepaid lease payments made less any lease incentives. The Company considered information available at the adoption date of Topic 842 to determine the incremental borrowing rate for leases in existence as of this date. The incremental borrowing rate used was the weighted average rate between secured and unsecured lending arrangements. Lease terms may include options to extend or terminate the lease when the Company is reasonably certain that the option will be exercised. Variable payments included in the lease agreement are expensed as incurred. Lease expense is recognized on a straight-line basis over the lease term. The Company elected to apply each of the practical expedients described in ASC Topic 842-10-65-1(f) which allow companies not to reassess: (i) whether any expired or existing agreements contain leases, (ii) the classification of any expired or existing leases, and (iii) the capitalization of initial direct costs for any existing leases. The Company also elected to apply the short-term lease measurement and recognition exemption in which ROU assets and lease liabilities are not recognized for short-term operating leases. A short-term operating lease is a lease that, at the commencement date, has a lease term of 12 months or less and does not include an option to purchase the underlying asset that the lessee is reasonably certain to exercise. The Company also elected not to separate non-lease components from lease components and instead to account for each separate lease component and the non-lease components associated with that lease component as a single lease component. Segment Information Segment Information The Company manages its operations as a single segment for the purposes of assessing performance and making operating decisions. The Company's operations are focused on oncology development activities. Research and Development Costs Research and Development Costs Costs incurred in connection with research and development activities are expensed as incurred. These costs include fees paid to consultants, vendors and various entities that perform certain research and testing on behalf of the Company. The Company has acquired rights to develop and commercialize product candidates. Upfront payments that relate to the acquisition of a new product compound, as well as pre-commercial milestone payments, are immediately expensed as acquired in-process research and development in the period in which they are incurred, provided that the new product compound did not also include processes or activities that would constitute a "business" as defined under U.S. GAAP, and the product candidate has not achieved regulatory approval for marketing and, absent obtaining such approval, has no established alternative future use. Milestone payments made to third parties subsequent to regulatory approval which meet the capitalization criteria would be capitalized as intangible assets and amortized over the estimated remaining useful life of the related product. If the conditions enabling capitalization of development costs as an asset have not yet been met, all development expenditures are recognized in profit or loss when incurred. Stock-based Compensation Stock-based Compensation The Company recognizes compensation cost for grants of employee stock options using a fair-value measurement method, that is recognized in operating results as compensation expense based on the fair value over the requisite service period of the awards. Forfeitures are recorded as they occur instead of estimating forfeitures that are expected to occur. The Company determines the fair value of stock-based awards that are based only on a service condition using the Black-Scholes option-pricing model which uses both historical and current market data to estimate fair value. The method incorporates various assumptions such as the risk-free interest rate, volatility, dividend yield, and expected life of the options. The Company determines the fair value of stock-based awards that are based on both a service condition and achievement of the first to occur of a market or performance condition using a Monte Carlo simulation. Income Taxes Income Taxes The Company accounts for income taxes using the asset and liability method, which requires the recognition of deferred tax assets and liabilities for the expected future tax consequences of events that have been recognized in the consolidated financial statements or in the Company's tax returns. The difference between the financial statement and tax basis of assets and liabilities is determined annually. Deferred income tax assets and liabilities are computed for those differences that have future tax consequences using the currently enacted tax laws and rates that apply to the years in which they are expected to affect taxable income. The Company assesses the likelihood that its deferred tax assets will be recovered from future taxable income and, to the extent it believes, based upon the weight of available evidence, that it is more likely than not that all or a portion of the deferred tax assets will not be realized, a valuation allowance is established through a charge to income tax expense. The Company uses a recognition threshold and measurement attribute for the financial statement recognition and measurement of a tax position taken, or expected to be taken, in a tax return. The provision for income taxes includes the effects of any resulting tax reserves, or unrecognized tax benefits, that are considered appropriate. The Company's policy is to record interest and penalties related to income taxes as part of the tax provision. Returns for tax years beginning with those filed for the period ended December 31, 2018 are open to federal and state tax examination. Accounting Pronouncements Not Yet Adopted Recent Accounting Pronouncements In November 2023, the FASB issued Topic 280 "Improvements to Reportable Segment Disclosures" which expands annual and interim disclosure requirements for reportable segments, primarily through enhanced disclosures about significant segment expenses. This guidance is effective for our annual periods beginning January 1, 2024, and for interim periods beginning January 1, 2025, with early adoption permitted. We are currently evaluating the potential effect that the updated standard will have on our financial statement disclosures. In December 2023, the FASB issued Topics 740 "Improvements to Income Tax Disclosures" to expand the disclosure requirements for income taxes, specifically related to the rate reconciliation and income taxes paid. This guidance is effective for our annual periods beginning January 1, 2025, with early adoption permitted. We are currently evaluating the potential effect that the updated standard will have on our financial statement disclosures. X - Definition Leases policy text block. + References No definition available. + Details Name: nuvb Leases Policy Text Block Namespace Prefix: nuvb Data Type: dtr-types:textBlockItem Type Balance Type: na Period Type: duration X - Definition Liquidity Policy Text Block. + References No definition available. + Details Name: nuvb Liquidity Policy Text Block Namespace Prefix: nuvb Data Type: dtr-types:textBlockItem Type Balance Type: na Period Type: duration X - Definition Revenue Recognition Policy [Policy Text Block] + References No definition available. + Details Name: nuvb Revenue Recognition Policy Policy Text Block Namespace Prefix: nuvb Data Type: dtr-types:textBlockItem Type Balance Type: na Period Type: duration X - Definition Unaudited Interim Financial Information policy. + References No definition available. + Details Name: nuvb Unaudited Interim Financial Information Policy Policy Text Block Namespace Prefix: nuvb Data Type: dtr-types:textBlockItem Type Balance Type: na Period Type: duration X - Definition Warrants policy text block. + References No definition available. + Details Name: nuvb Warrants Policy Text Block Namespace Prefix: nuvb Data Type: dtr-types:textBlockItem Type Balance Type: na Period Type: duration X - References No definition available. + Details Name: us-gaap Accounting Policies Abstract Namespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - Definition Disclosure of accounting policy for basis of accounting, or basis of presentation, used to prepare the financial statements (for example, US Generally Accepted Accounting Principles, Other Comprehensive Basis of Accounting, IFRS). + References No definition available. + Details Name: us-gaap Basis Of Accounting Policy Policy Text Block Namespace Prefix: us-gaap Data Type: dtr-types:textBlockItem Type Balance Type: na Period Type: duration X - Definition Disclosure of accounting policy for cash and cash equivalents, including the policy for determining which items are treated as cash equivalents. Other information that may be disclosed includes (1) the nature of any restrictions on the entity's use of its cash and cash equivalents, (2) whether the entity's cash and cash equivalents are insured or expose the entity to credit risk, (3) the classification of any negative balance accounts (overdrafts), and (4) the carrying basis of cash equivalents (for example, at cost) and whether the carrying amount of cash equivalents approximates fair value. + References Reference 1:

Codification -Section S99 -Paragraph 2 -Subparagraph (SAB Topic 5.CC) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480091/360-10-S99-2Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Section 05 -Paragraph 4 -SubTopic 10 -Topic 360 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482338/360-10-05-4 + Details Name: us-gaap ImpairmentOrDisposalOfLongLivedAssetsPolicyTextBlock Namespace Prefix: us-gaap_ Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - DefinitionDisclosure of accounting policy for income taxes, which may include its accounting policies for recognizing and measuring deferred tax assets and liabilities and related valuation allowances, recognizing investment tax credits, operating loss carryforwards, tax credit carryforwards, and other carryforwards, methodologies for determining its effective income tax rate and the characterization of interest and penalties in the financial statements. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 740 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 20 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482685/740-10-50-20Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 740 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 19 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482685/740-10-50-19Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 740 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 25 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482525/740-10-45-25Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-03(h)(1)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479886/946-10-S99-3Reference 5: http://www.xbrl.org/2003/role/disclosureRef -Topic 740 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 17 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482685/740-10-50-17Reference 6: http://www.xbrl.org/2003/role/disclosureRef -Topic 740 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 9 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482685/740-10-50-9Reference 7: http://www.xbrl.org/2003/role/disclosureRef -Topic 740 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482525/740-10-45-28Reference 8: http://www.xbrl.org/2003/role/disclosureRef -Topic 220 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Publisher FASB -URI https://asc.fasb.org/1943274/2147477250/946-10-50-1Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-03(d)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479886/946-10-S99-3Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-03(f)(1)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479886/946-10-S99-3Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-03(f)(2)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479886/946-10-S99-3Reference 5: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-03(f)(3)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479886/946-10-S99-3Reference 6: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 320 -Name Accounting Standards Codification -Section S99 -Paragraph 12 -Subparagraph (i) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477271/946-320-S99-12Reference 7: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 320 -Name Accounting Standards Codification -Section S99 -Paragraph 19 -Subparagraph (2) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477271/946-320-S99-19 + Details Name: us-gaap InvestmentPolicyTextBlock Namespace Prefix: us-gaap_ Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - DefinitionDisclosure of accounting policy pertaining to new accounting pronouncements that may impact the entity's financial reporting. Includes, but is not limited to, quantification of the expected or actual impact. + ReferencesNo definition available. + Details Name: us-gaap NewAccountingPronouncementsPolicyTextBlock Namespace Prefix: us-gaap_ Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - DefinitionDisclosure of accounting policy for long-lived, physical asset used in normal conduct of business and not intended for resale. Includes, but is not limited to, work of art, historical treasure, and similar asset classified as collections. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Section 50 -Paragraph 1 -SubTopic 10 -Topic 360 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482099/360-10-50-1Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(8)(a)) -Publisher FASB -URI https://asc.fasb.org/1943274/214747877/944-210-S99-1Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Name Accounting Standards Codification -Section 50 -Paragraph 6 -SubTopic 360 -Topic 958 -Publisher FASB -URI https://asc.fasb.org/1943274/2147477798/958-360-50-6Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (d) -SubTopic 360 -Topic 958 -Publisher FASB -URI https://asc.fasb.org/1943274/2147477798/958-360-50-1 + Details Name: us-gaap PropertyPlantAndEquipmentPolicyTextBlock Namespace Prefix: us-gaap_ Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - DefinitionDisclosure of accounting policy for costs it has incurred (1) in a planned search or critical investigation aimed at discovery of new knowledge with the hope that such knowledge will be useful in developing a new product or service, a new process or technique, or in bringing about a significant improvement to an existing product or process; or (2) to translate research findings or other knowledge into a plan or design for a new product or process or for a significant improvement to an existing product or process. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 730 -SubTopic 10 -Name Accounting Standards Codification -Section 05 -Paragraph 1 -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483044/730-10-05-1 + Details Name: us-gaap ResearchAndDevelopmentExpensePolicy Namespace Prefix: us-gaap_ Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - DefinitionDisclosure of accounting policy for revenue from contract with customer. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 17 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-17Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 19 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-19Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 18 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-18Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 18 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-18Reference 5: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 20 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-20Reference 6: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 20 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-20Reference 7: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 20 -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-20Reference 8: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 20 -Subparagraph (d) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-20Reference 9: http://www.xbrl.org/2003/role/exampleRef -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph (e) -SubTopic 10 -Topic 235 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483426/235-10-50-4Reference 10: http://www.xbrl.org/2003/role/disclosureRef -Name Accounting Standards Codification -Topic 606 -Publisher FASB -URI https://asc.fasb.org/606/tableOfContent + Details Name: us-gaap RevenueFromContractWithCustomerPolicyTextBlock Namespace Prefix: us-gaap_ Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - DefinitionDisclosure of accounting policy for segment reporting. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 41 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-41Reference 2: http://www.xbrl.org/2003/role/exampleRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 47 -Subparagraph (bb) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482785/280-10-55-47Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 29 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-29Reference 4: http://www.xbrl.org/2003/role/exampleRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 54 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482785/280-10-55-54Reference 5: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 36 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-36Reference 6: http://www.xbrl.org/2003/role/exampleRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 47 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482785/280-10-55-47Reference 7: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 29 -Subparagraph (d) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-29Reference 8: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 29 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-29Reference 9: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 29 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-29Reference 10: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 29 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-29Reference 11: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 29 -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-29Reference 12: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 29 -Subparagraph (e) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-29 + Details Name: us-gaap SegmentReportingPolicyTextBlock Namespace Prefix: us-gaap_ Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - DefinitionDisclosure of accounting policy for award under share-based payment arrangement. Includes, but is not limited to, methodology and assumption used in measuring cost. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (f)(1) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (f)(2)(i) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (f)(2)(ii) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (f)(2)(v) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2Reference 5: http://www.xbrl.org/2003/role/disclosureRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SAB Topic 14.C.Q3) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479830/718-10-S99-1Reference 6: http://www.xbrl.org/2003/role/disclosureRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SAB Topic 14.D.1.Q5) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479830/718-10-S99-1Reference 7: http://www.xbrl.org/2003/role/disclosureRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SAB Topic 14.D.3.Q2) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479830/718-10-S99-1Reference 8: http://www.xbrl.org/2003/role/disclosureRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SAB Topic 14.D.2.Q6) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479830/718-10-S99-1Reference 9: http://www.xbrl.org/2003/role/disclosureRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Publisher FASB -URI https://asc.fasb.org/718/tableOfContentReference 10: http://www.xbrl.org/2003/role/disclosureRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2Reference 11: http://www.xbrl.org/2003/role/disclosureRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (f)(2) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2 + Details Name: us-gaap ShareBasedCompensationOptionAndIncentivePlansPolicy Namespace Prefix: us-gaap_ Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - DefinitionTabular disclosure of the nature of the unusual risk or uncertainty, if estimable, such as the threat of expropriation of its assets by a foreign government, rapid technological obsolescence in the industry, risk of natural disaster from earthquake or weather events, and availability of or continuation of a labor force at a reasonable cost. + ReferencesNo definition available. + Details Name: us-gaap UnusualRisksAndUncertaintiesTextBlock Namespace Prefix: us-gaap_ Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - DefinitionDisclosure of accounting policy for the use of estimates in the preparation of financial statements in conformity with generally accepted accounting principles. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Topic 275 -SubTopic 10 -Section 50 -Paragraph 9 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482861/275-10-50-9Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Topic 275 -SubTopic 10 -Section 50 -Paragraph 4 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482861/275-10-50-4Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (b) -SubTopic 10 -Topic 275 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482861/275-10-50-1Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (c) -SubTopic 10 -Topic 275 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482861/275-10-50-1Reference 5: http://www.xbrl.org/2003/role/disclosureRef -Name Accounting Standards Codification -Section 50 -Paragraph 11 -SubTopic 10 -Topic 275 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482861/275-10-50-11Reference 6: http://www.xbrl.org/2003/role/disclosureRef -Name Accounting Standards Codification -Section 50 -Paragraph 12 -SubTopic 10 -Topic 275 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482861/275-10-50-12Reference 7: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Topic 275 -SubTopic 10 -Section 50 -Paragraph 8 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482861/275-10-50-8 + Details Name: us-gaap UseOfEstimates Namespace Prefix: us-gaap_ Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration XML 29 R22.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Significant Accounting Policies (Tables) 6 Months Ended Jun. 30, 2024 Accounting Policies [Abstract] A Schedule of Concentration of Risk Concentration of customers Concentration of customers Substantially all revenues were derived from customers located in China and Japan. For the three and six months ended June 30, 2024, the revenues derived from customers located in China and Japan were \$1.1 million and \$0.3 million, respectively. There was no revenue in 2023 since the Acquisition of AnHeart completed in the second quarter of 2024.Below customers represent more than 10% of the Company's net revenue for the three and six months ended June 30, 2024. A A Three Months Ended June 30, A A Six Months Ended June 30, A A 2024 A A 2024 A Customer A A A 1,096 A A A 1,096 A Customer B A A 339 A A A 339 A A A A A A Below customer represents more than 10% of the Company's balances of accounts receivable as of June 30, 2024. A A June 30, 2024 A Customer B A A 117 A A A A X - ReferencesNo definition available. + Details Name: us-gaap AccountingPoliciesAbstract Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionTabular disclosure of the nature of a concentration, a benchmark to which it is compared, and the percentage that the risk is to the benchmark. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 825 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 21 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482907/825-10-50-21Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Topic 825 -SubTopic 10 -Section 50 -Paragraph 20 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482907/825-10-50-20Reference 3: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Topic 275 -SubTopic 10 -Section 50 -Paragraph 18 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482861/275-10-50-18Reference 4: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Topic 275 -SubTopic 10 -Section 50 -Paragraph 20 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482861/275-10-50-20Reference 5: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Topic 275 -SubTopic 10 -Section 50 -Paragraph 16 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482861/275-10-50-16Reference 6: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -

[Topic 275 - SubTopic 10 - Section 50 - Paragraph 8](#) Publisher FASB -URI https://asc.fasb.org/1943274/2147482861/275-10-50-2+ Details Name: us-gaap_ScheduleOfConcentrationOfRiskByRiskFactorTextBlock Namespace Prefix: us-gaap Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration XML 30 R23.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Acquisition (Tables) 6 Months Ended Jun. 30, 2024 Business Combinations [Abstract] A Schedule of Final Allocable Purchase Price Consists The final allocable purchase price consists of the following (in thousands): Class A Common Stock (1) \$ 89,297 Series A Non-Voting Convertible Preferred Stock (2) \$ 274,938 Consideration Warrants (3) \$ 764 Assumed share-based awards (4) \$ 24,818 Acquisition costs \$ 4,734 Total allocable purchase price \$ 397,251 A A A A
27,646,255 shares issued at \$3.23 per share as of the Acquisition Date. (2) 851,202 shares issued at as-converted value of \$323 per share.(3) 2,893,731 warrants issued at the estimated fair value of \$0.2641 per warrant.(4) The portion recorded in consideration transferred represents the fair value of replacement awards (Assumed Options & Assumed RSUs) attributable to precombination services. X - ReferencesNo definition available. + Details Name: us-gaap_BusinessCombinationsAbstract Namespace Prefix: us-gaap Data Type: xbrl:stringItemType Balance Type: na Period Type: duration X - DefinitionTabular disclosure of a material business combination completed during the period, including background, timing, and recognized assets and liabilities. This table does not include leveraged buyouts. + ReferenceReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Topic 805 -SubTopic 10 -Section 50 -Paragraph 3 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479328/805-10-50-3Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Topic 805 -SubTopic 10 -Section 50 -Paragraph 2 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479328/805-10-50-2+ Details Name: us-gaap_ScheduleOfBusinessAcquisitionTextBlock Namespace Prefix: us-gaap Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration XML 31 R24.htm IDEA: XBRL DOCUMENT V3.24.2.u1 Fair Value Measurements and Marketable Securities Available-for-Sale (Tables) 6 Months Ended Jun. 30, 2024 Fair Value Disclosures [Abstract] A Schedule of Marketable securities Measured at Fair Value on Recurring Basis There have not been any transfers between the levels during the periods.A A June 30, 2024 A A Total A A Level 1A A Level 2A A Level 3A A (In thousands) A Financial assets: A A A A Cash equivalents: A A A A Money market funds A \$ 7,492 A A \$ 7,492 A A \$ æ” A A \$ æ” U.S. government securities A 9,926 A A æ” A A 9,926 A A æ” Corporate bonds A 2,138 A A æ” A A 2,138 A A æ” A A 19,556 A A \$ 7,492 A A 12,064 A A æ” A A Marketable securities: A A A A Certificate of deposits A 21,694 A A æ” A A 21,694 A A æ” Commercial paper A 36,555 A A æ” A A 36,555 A A æ” A A U.S. government and government agency securities A 408,996 A A æ” A A 408,996 A A æ” Corporate bonds A 75,639 A A æ” A A 75,639 A A æ” A A 542,884 A A æ” A A 542,884 A A æ” Total financial assets: A \$ 562,440 A \$ 7,492 A \$ 554,948 A A \$ æ” A A A A Financial liabilities: A A A A Warrants A \$ 1,441 A A \$ 1,363 A A \$ æ” A A 78 A A December 31, 2023 A A Total A A Level 1A A Level 2A A Level 3A A (In thousands) A Financial assets: A A A A Cash equivalents: A A A A Money market funds A \$ 20,574 A A \$ 20,574 A A \$ æ” A A \$ æ” U.S. government and government agency securities A 4,988 A A æ” A A 4,988 A A æ” Corporate bonds A 947 A A æ” A A 947 A A æ” A A 26,509 A A \$ 20,574 A A 5,935 A A æ” Marketable securities: A A A A Certificate of deposits A 10,176 A A A A æ” A A 10,176 A A æ” Commercial paper A 23,628 A A æ” A A 23,628 A A æ” U.S. government and government agency securities A 453,175 A A æ” A A 453,175 A A æ” Corporate bonds A 81,585 A A æ” A A 81,585 A A æ” A A 568,564 A A æ” A A 568,564 A A æ” Total financial assets: A \$ 595,073 A A \$ 20,574 A A \$ 574,499 A A \$ æ” A A A A Financial liabilities: A A A A Warrants A \$ 353 A A \$ 338 A A \$ æ” A A \$ 15 A Schedule of Cost, Aggregate Fair Value and Unrealized Gains of Marketable Securities Available-for-sale The following table provides the amortized cost, aggregate fair value, and unrealized gains (losses) of marketable securities as of June 30, 2024 and December 31, 2023:A A June 30, 2024 A A AmortizedCost A A UnrealizedGains A A Unrealized Losses A A Fair Value A A (In thousands) A Cash equivalents: A A A A Money market funds A \$ 7,492 A A \$ æ” A A \$ 7,492 A A U.S. government securities A 9,926 A A æ” A A 9,926 A A æ” Corporate bonds A 2,138 A A æ” A A 2,138 A A æ” A A 19,556 A A æ” A A 19,556 A A æ” Certificate of deposits A 21,689 A A 8 A A (3) A A 21,694 A A Commercial paper A 36,580 A A 4 A A (29) A A 36,555 A A U.S. government and government agency securities A 410,234 A A 10 A A (1,248) A A 408,996 A A Corporate bonds A 75,795 A A 42 A A (198) A A 75,639 A A A 544,298 A A 64 A A (1,478) A A 542,884 A A \$ 563,854 A A \$ 64 A A (\$ 1,478) A A \$ 562,440 A A A A December 31, 2023 A A AmortizedCost A A UnrealizedGains A A Unrealized Losses A A Fair Value A A (In thousands) A Cash equivalents: A A A A Money market funds A \$ 20,574 A A \$ æ” A A \$ 20,574 A A U.S. government and government agency securities A 4,987 A A 1 A A æ” A A 4,988 A A Corporate bonds A 947 A A æ” A A 947 A A æ” A A 26,508 A A 1 A A æ” A A 26,509 A A Marketable securities: A A A A A A Certificate of deposits A 10,171 A A 7 A A (2) A A 10,176 A A Commercial paper A 23,609 A A 22 A A (3) A A 23,628 A A U.S. government and government agency securities A 452,813 A A 1,095 A A (733) A A 453,175 A A Corporate bonds A 81,791 A A 68 A A (274) A A 81,585 A A A A 568,384 A A 1,192 A A (1,012) A A 568,564 A A \$ 594,892 A A \$ 1,193 A A (\$ 1,012) A A \$ 595,073 A A Schedule of Net Realized Gain (Loss) on Marketable Securities For the three and six months ended June 30, 2024 and 2023, there were no realized gains or losses on equity securities. The activity related to the net gains (losses) on marketable securities included in other income (expense) on the consolidated statements of operations and comprehensive loss were as follows (in thousands):A A Three Months Ended June 30, A A Six Months Ended June 30, A A 2024 A A 2023 A A 2024 A A 2023 A A Net realized gains (losses) on available-for-sale securities were as follows: A A A A A A Realized gains from sales of available-for-sale securities A \$ 1 A A \$ 12 A A \$ 1 A A \$ 12 A A Realized losses from sales of available-for-sale securities A (8) A A (11) A A (7) A A (207) A A Total losses on marketable securities A (7) A A (99) A A (6) A A (195) A A A A A A Schedule of Marketable Securities With Continuous Unrealized Losses The following tables provide marketable securities with continuous unrealized losses for less than 12 months and 12 months or greater and the related fair values as of June 30, 2024 and December 31, 2023 were as follows: A A June 30, 2024 A A Less than 12 Months A A 12 Months or Greater A A Total A A Total A A Fair Value A A Unrealized Losses A A Fair Value A A Unrealized Losses A A Fair Value A A Unrealized Losses A A (In thousands) A Certificate of deposits A \$ 9,940 A A \$ (3) A A \$ æ” A A \$ 9,940 A A \$ (3) Commercial paper A 26,185 A A (29) A A æ” A A 26,185 A A (29) A A U.S. government and government agency securities A 270,609 A A (628) A A 115,683 A A (620) A A 386,292 A A (1,248) A A Corporate bonds A 46,561 A A (70) A A 11,582 A A (128) A A 58,143 A A (198) A A 353,295 A A (730) A A 127,265 A A (748) A A \$ 480,560 A A \$ (1,478) A A December 31, 2023 A A Less than 12 Months A A 12 Months or Greater A A Total A A Total A A Fair Value A A Unrealized Losses A A Fair Value A A Unrealized Losses A A Fair Value A A Unrealized Losses A A (In thousands) A Certificate of deposits A \$ 5,693 A A \$ (2) A A \$ æ” A A \$ 5,693 A A \$ (2) Commercial paper A 9,101 A A (3) A A æ” A A 9,101 A A (3) A A U.S. government and government agency securities A 199,552 A A (694) A A 24,761 A A (38) A A 224,313 A A (732) A A Corporate bonds A 21,844 A A (28) A A 26,484 A A (246) A A 48,328 A A (274) A A \$ 236,190 A A \$ (727) A A \$ 51,245 A A (284) A A \$ 287,435 A A \$ (1,011) A A Schedule Of Available for sale Securities Maturity On Fair Value Maturity information based on fair value is as follows as of June 30, 2024:A A Within one year A A After one yearthrough five years A A Total A A (In thousands) A Certificate of deposits A \$ 21,069 A A \$ 625 A A \$ 21,694 A A Commercial paper A 36,555 A A æ” A A 36,555 A A U.S. government and government agency securities A 283,225 A A 125,771 A A 408,996 A A Corporate bonds A 55,284 A A 20,355 A A 75,639 A A \$ 396,133 A A \$ 146,751 A A \$ 542,884 A A X -DefinitionSchedule Of Available for sale Securities Classified By Contractual Maturity Date Table Text Block + ReferencesNo definition available. + Details Name: nuvb_ScheduleOfAvailableForSaleSecuritiesClassifiedByContractualMaturityDateTableTextBlock Namespace Prefix: nuvb Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - DefinitionTabular disclosure of fair value of investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available for sale), in unrealized loss position, without allowance for credit loss. Includes beneficial interest in securitized financial asset. + ReferenceReference 1:
<http://www.xbrl.org/2003/role/exampleRef>-Topic 326 -SubTopic 30 -Name Accounting Standards Codification -Section 55 -Paragraph 8 -Publisher FASB -URI
<https://asc.fasb.org/1943274/2147479081/326-30-55-8Reference 2: http://fasb.org/us-gaap/role/ref/otherTransitionRef>-Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 6 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-6Reference 3: <http://www.xbrl.org/2009/role/commonPracticeRef>-Topic 326 -SubTopic 30 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479106/326-30-50-4+ Details Name: us-gaap_DebtSecuritiesAvailableFor

Period Type: duration XML 36 R29.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Stock-Based Compensation (Tables) 6 Months Ended Jun. 30, 2024 Share-Based Payment Arrangement [Abstract] Â Summary of Stock-Based Compensation Expense The stock-based compensation expense included in the Companyâ€™s condensed statement of operations and comprehensive loss for the three and six months ended June 30, 2024 and 2023 is as follows (in thousands): Â Â Â Three Months Ended June 30, Â Â Â Six Months Ended June 30, Â Â Â 2024 Â Â 2023 Â Â 2024 Â Â 2023 Â Research and development Â \$,4190 Â \$,2,420 Â \$,6,482 Â \$,4,431 Â General and administrative Â \$,5,379 Â \$,3,103 Â \$,8,128 Â \$,5,929 Â \$,9,569 Â \$,5,523 Â Â \$,14,610 Â Â \$,10,360 Â Â Summary of Stock Option Activity under 2019 Plan and 2021 Plan The following table summarizes stock option activity under the 2019 Plan, 2021 Plan and assumed options in the AnHeart Acquisition for the six months ended June 30, 2024. Â Â Shares Â Outstanding at December 31, 2023 Â Â 30,649,239 Â Options assumed for the Acquisition of AnHeart Â Â 13,742,239 Â Granted Â Â 11,322,679 Â Exercised Â Â (1,798,190) Forfeited or canceled Â Â (1,223,019) Outstanding at June 30, 2024 Â Â 52,692,948 Â Exercisable at June 30, 2024 Â Â 23,909,107 Â Â Summary of Restricted Stock Units Activity The following table summarizes the activity for the restricted stock units assumed in the AnHeart Acquisition for the six months ended June 30, 2024. Â Â Â Restricted Stock Â Â Â Units Â Non-vested at December 31, 2023 Â Â â€” Â RSUs assumed for the Acquisition of AnHeart Â Â 2,201,694 Â Granted Â Â â€” Â Vested Â Â (40,342) Forfeited or canceled Â Â â€” Â Non-vested at June 30, 2024 Â Â 2,161,352 Â Â Â Â Â X - ReferencesNo definition available. + Details Name: us-gaap DisclosureOfCompensationRelatedCostsSharebasedPaymentsAbstract Namespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionTabular disclosure of allocation of amount expensed and capitalized for award under share-based payment arrangement to statement of income or comprehensive income and statement of financial position. Includes, but is not limited to, corresponding line item in financial statement. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Topic 718 -SubTopic 10 -Section 50 -Paragraph 2 -Subparagraph (h)(1) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2 + Details Name: us-gaap ScheduleOfEmployeeServiceShareBasedCompensationAllocationOfRecognizedPeriodCostsTextBlock Namespace Prefix: us-gaap Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - DefinitionTabular disclosure of the number and weighted-average grant date fair value for restricted stock units that were outstanding at the beginning and end of the year, and the number of restricted stock units that were granted, vested, or forfeited during the year. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2 + Details Name: us-gaap ScheduleOfShareBasedCompensationRestrictedStockUnitsAwardActivityTableTextBlock Namespace Prefix: us-gaap Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - DefinitionTabular disclosure for stock option plans. Includes, but is not limited to, outstanding awards at beginning and end of year, grants, exercises, forfeitures, and weighted-average grant date fair value. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (d) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2Reference 3: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (e) -SubTopic 10 -Topic 718 -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2 + Details Name: us-gaap ScheduleOfShareBasedCompensationStockOptionsActivityTableTextBlock Namespace Prefix: us-gaap Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration XML 37 R30.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Warrants (Tables) 6 Months Ended Jun. 30, 2024 Warrant Disclosure [Abstract] Â Schedule of Quantitative Information Regarding Level 3 Fair Value Measurements The following table provides quantitative assumptions regarding Level 3 fair value measurements: Â Â Â June 30, 2024 Â Â December 31, 2023 Â Stock price Â \$ 2.92 Â Â \$ 1.51 Â Strike price Â \$ 11.50 Â Â \$ 11.50 Â Term (in years) Â 1.6 Â Â 2.1 Â Volatility Â 98.1 % Â 82.7 % Risk-free rate Â 4.9 % Â 4.1 % Dividend yield Â 0.0 % Â 0.0 % Schedule of Fair Values for Outstanding Warrants The Company determined the following fair values for the outstanding warrants (in thousands): Â Â Â June 30, 2024 Â Public Warrants Â \$ 767 Â Private Placement Warrants Â 78 Â Forward Purchase Warrants Â 133 Â Consideration Warrants Â 463 Â Total Â \$ 1,441 Â Summary of Changes in liabilities classified in Level 3 of the fair value hierarchy The following presents changes in liabilities classified in Level 3 of the fair value hierarchy for the six months ended June 30, 2024 (in thousands): Â Â Six Months Ended Â Â June 30, 2024 Â Beginning balance Â \$ 15 Â Change in fair value of Private Warrants liability recognized in earnings Â 63 Â Ending balance Â \$ 78 Â Â Â Â X - DefinitionWarrant disclosure. + ReferencesNo definition available. + Details Name: nuvb WarrantDisclosureAbstract Namespace Prefix: nuvb Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionTabular disclosure of input and valuation technique used to measure fair value and change in valuation approach and technique for each separate class of asset and liability measured on recurring and nonrecurring basis. + ReferencesReference 1: http://www.xbrl.org/2003/role/exampleRef -Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 103 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482078/820-10-55-103Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (bbb)(2) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (bbb)(1) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2 + Details Name: us-gaap FairValueAssetsAndLiabilitiesMeasuredOnRecurringAndNonrecurringBasisValuationTechniquesTableTextBlock Namespace Prefix: us-gaap Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - DefinitionTabular disclosure of liabilities, including [financial] instruments measured at fair value that are classified in stockholders' equity, if any, by class that are measured at fair value on a recurring basis. The disclosures contemplated herein include the fair value measurements at the reporting date by the level within the fair value hierarchy in which the fair value measurements in their entirety fall, segregating fair value measurements using quoted prices in active markets for identical assets (Level 1), significant other observable inputs (Level 2), and significant unobservable inputs (Level 3). Where the quoted price in an active market for the identical liability is not available, the Level 1 input is the quoted price of an identical liability when traded as an asset. + ReferencesReference 1: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2Reference 2: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2 + Details Name: us-gaap FairValueLiabilitiesMeasuredOnRecurringBasisTextBlock Namespace Prefix: us-gaap Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration X - DefinitionTabular disclosure of warrants or rights issued. Warrants and rights outstanding are derivative securities that give the holder the right to purchase securities (usually equity) from the issuer at a specific price within a certain time frame. Warrants are often included in a new debt issue to entice investors by a higher return potential. The main difference between warrants and call options is that warrants are issued and guaranteed by the company, whereas options are exchange instruments and are not issued by the company. Also, the lifetime of a warrant is often measured in years, while the lifetime of a typical option is measured in months. Disclose the title of issue of securities called for by warrants and rights outstanding, the aggregate amount of securities called for by warrants and rights outstanding, the date from which the warrants or rights are exercisable, and the price at which the warrant or right is exercisable. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Topic 718 -SubTopic 10 -Section 50 -Paragraph 1 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-1 + Details Name: us-gaap ScheduleOfStockholdersEquityNoteWarrantsOrRightsTextBlock Namespace Prefix: us-gaap Data Type: dtr-types:textBlockItemType Balance Type: na Period Type: duration XML 38 R31.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Significant Accounting Policies - Additional Information (Details) - USD (\$) \$ / shares in Units, \$ in Thousands 3 Months Ended 6 Months Ended 12 Months Ended Jun. 30, 2024 Jun. 30, 2024 Jun. 30, 2023 Dec. 31, 2023 Accumulated deficit \$ (820,088) \$ (820,088) \$ (342,804) Net cash used in operating activities Â \$ (53,026) \$ (36,815) Â Sale of Stock, Description of Transaction Â the exercise of the Warrants may be settled in cash upon the occurrence of a tender offer or exchange that involves more than 50% of the outstanding shares of the Companyâ€™s Common Stock. Â Â Cash and cash equivalents and marketable securities 577,200 \$ 577,200 Â Â AnHeart [Member] Â Â Â Â Revenue from customer 0 Â Â \$ 0 China and Japan [Member] Â Â Â Â Revenue from customer \$,100 \$ 300 Â Â Revenue Benchmark [Member] | Customer Concentration Risk [Member] Â Â Â Â Concentration Risk, Percentage 10.00% 10.00% Â Â Accounts Receivable [Member] | Customer Concentration Risk [Member] Â Â Â Â Concentration Risk, Percentage 10.00% 10.00% Â Â Class A Common Stock [Member] Â Â Â Â Warrants to purchase common stock outstanding 5,787,472 5,787,472 Â Â Warrants [Member] Â Â Â Â Warrants to purchase common stock outstanding 5,787,472 5,787,472 Â Â Class of warrant or right, exercise price of warrants or rights \$ 11.5 \$ 11.5 Â Â Public Warrants [Member] Â Â Â Â Warrants to purchase common stock outstanding 4,791,639 4,791,639 Â Â Private Placement Warrants [Member] Â Â Â Â Warrants to purchase common stock outstanding 162,500 162,500 Â Â Forward Purchase Warrants [Member] Â Â Â Â Warrants to purchase common stock outstanding 833,333 833,333 Â Â Consideration Warrants [Member] | Class A Common Stock [Member] Â Â Â Â Warrants to purchase common stock outstanding 2,893,731 2,893,731 Â Â Class of warrant or right, exercise price of warrants or rights \$ 11.5 \$ 11.5 Â Â Fair value of warrants, trading price \$ 0.26 \$ 0.26 Â Â Computer Equipment [Member] Â Â Â Â Property and equipment, estimated useful life 5 years 5 years Â Â Furniture and Equipment [Member] Â Â Â Â Property and equipment, estimated useful life 7 years 7 years Â Â X - DefinitionCash and cash equivalents and marketable securities. + ReferencesNo definition available. + Details Name: nuvb CashAndCashEquivalentsAndMarketableSecurities Namespace Prefix: nuvb Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionConcentration Risk Threshold Percentage + ReferencesNo definition available. + Details Name: nuvb ConcentrationRiskThresholdPercentage Namespace Prefix: nuvb Data Type: dtr-types:percentItemType Balance Type: na Period Type: duration X - DefinitionFair Value of Warrants, Trading Price + ReferencesNo definition available. + Details Name: nuvb FairValueOfWarrantsTradingPrice Namespace Prefix: nuvb Data Type: dtr-types:perShareItemType Balance Type: na Period Type: instant X - DefinitionExercise price per share or per unit of warrants or rights outstanding. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481112/505-10-50-3 + Details Name: us-gaap ClassOfWarrantOrRightExercisePriceOfWarrantsOrRights1 Namespace Prefix: us-gaap Data Type: dtr-types:perShareItemType Balance Type: na Period Type: instant X - DefinitionNumber of securities into which the class of warrant or right may be converted. For example, but not limited to, 500,000 warrants may be converted into 1,000,000 shares. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481112/505-10-50-3 + Details Name: us-gaap ClassOfWarrantOrRightNumberOfSecuritiesCalledByWarrantsOrRights Namespace Prefix: us-gaap Data Type: xbrli:sharesItemType Balance Type: na Period Type: instant X - DefinitionAmount of cash inflow (outflow) from operating activities, including discontinued operations. Operating activity cash flows include transactions, adjustments, and changes in value not defined as investing or financing activities. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 230 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 28 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482740/230-10-45-28Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 230 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 24 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482740/230-10-45-24Reference 3: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 230 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 25 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482740/230-10-45-25 + Details Name: us-gaap NetCashProvidedByUsedInOperatingActivities Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: na Period Type: duration X - DefinitionUseful life of long lived, physical assets used in the normal conduct of business and not intended for resale, in 'PnYnMnDnTnHnMns' format, for example, 'P1Y5M13D' represents the reported fact of one year, five months, and thirteen days. Examples include, but not limited to, land, buildings, machinery and equipment, furniture and fixtures, and computer equipment. + ReferencesNo definition available. + Details Name: us-gaap PropertyPlantAndEquipmentUsefulLife Namespace Prefix: us-gaap Data Type: xbrli:durationItemType Balance Type: na Period Type: instant X - DefinitionAmount of accumulated undistributed earnings (deficit). + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(30)(a)(3)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 2: http://www.xbrl.org/2003/role/exampleRef -Topic 852 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 10 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481372/852-10-55-10Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 2 -Subparagraph (g)(2)(i) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480016/944-40-65-2Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 40 -Name Accounting Standards Codification -Section 65 -Paragraph 2 -Subparagraph (h)(2) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480016/944-40-65-2Reference 5: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 11 -Publisher FASB -URI https://asc.fasb.org/1943274/2147480990/946-20-50-11Reference 6: http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(23)(4)(4)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478777/944-210-S99-1Reference 7: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.6-04(17)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479170/946-210-S99-1Reference 8: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.3-04) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480008/505-10-S99-1 + Details Name: us-gaap RetainedEarningsAccumulatedDeficit Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionAmount, including tax collected from customer, of revenue from satisfaction of performance obligation by transferring promised good or service to customer. Tax collected from customer is tax assessed by governmental authority that is both imposed on and concurrent with specific revenue-producing transaction, including, but not limited to, sales, use, value-added and excise. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 41 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-41Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 270 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (i) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482964/270-10-50-1Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (ee) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference 4: http://fasb.org/us-gaap/role/ref/otherTransitionRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference 5: http://fasb.org/us-gaap/role/ref/otherTransitionRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference 6: http://www.xbrl.org/2003/role/disclosureRef -Topic 924 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SAB Topic 11.L) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479941/924-10-S99-1Reference 7: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 5 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-5Reference 8: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 30 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-30Reference 9: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 42 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-42Reference 10: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22Reference 11: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 40 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-40Reference 12: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22Reference 13: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic

10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-4 + Details Name: us-gaap RevenueFromContractWithCustomerIncludingAssessedTax Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionDescription of stock transaction which may include details of the offering (IPO, private placement), a description of the stock sold, percentage of subsidiary's or equity investee's stock sold, a description of the investors and whether the stock was issued in a business combination. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 810 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 23 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481231/810-10-45-23Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Topic 810 -SubTopic 10 -Section 50 -Paragraph 1A -Publisher FASB -URI https://asc.fasb.org/1943274/2147481203/810-10-50-1AReference 3: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 50 -Paragraph 1B -SubTopic 10 -Topic 810 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481203/810-10-50-1B + Details Name: us-gaap SaleOfStockDescriptionOfTransaction Namespace Prefix: us-gaap_ Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - Details Name: us-gaap BusinessAcquisitionAxis=nuvb AnheartMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: srt StatementGeographicalAxis=nuvb ChinaAndJapanMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap ConcentrationRiskByBenchmarkAxis=us-gaap SalesRevenueNetMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap ConcentrationRiskByTypeAxis=us-gaap CustomerConcentrationRiskMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap ConcentrationRiskByBenchmarkAxis=us-gaap AccountsReceivableMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap StatementClassOfStockAxis=us-gaap CommonClassAMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap ClassOfWarrantOrRightAxis=us-gaap WarrantMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap ClassOfWarrantOrRightAxis=nuvb PrivatePlacementWarrantsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap ClassOfWarrantOrRightAxis=nuvb ForwardPurchaseWarrantsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap ClassOfWarrantOrRightAxis=nuvb ConsiderationWarrantsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap PropertyPlantAndEquipmentByTypeAxis=us-gaap ComputerEquipmentMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap PropertyPlantAndEquipmentByTypeAxis=us-gaap FurnitureAndFixturesMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 39 R32.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Significant Accounting Policies - Schedule of Concentration of Credit Risk Net Revenue (Details) - USD (\$) \$ in Thousands 3 Months Ended 6 Months Ended Jun. 30, 2024 Jun. 30, 2024 Customer A [Member] A A Concentration Risk [Line Items] A A Revenue from customer \$ 1,096 \$ 1,096 Customer B [Member] A A Concentration Risk [Line Items] A A Revenue from customer \$ 339 \$ 339 X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 954 -SubTopic 310 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Publisher FASB -URI https://asc.fasb.org/1943274/2147478785/954-310-50-2 + Details Name: us-gaap ConcentrationRiskLineItems Namespace Prefix: us-gaap_ Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - DefinitionAmount, including tax collected from customer, of revenue from satisfaction of performance obligation by transferring promised good or service to customer. Tax collected from customer is tax assessed by governmental authority that is both imposed on and concurrent with specific revenue-producing transaction, including, but not limited to, sales, use, value-added and excise. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 41 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-41Reference 2: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 270 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (i) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482964/270-10-50-1Reference 3: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (ee) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference 4: <http://fasb.org/us-gaap/role/ref/otherTransitionRef> -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference 5: <http://fasb.org/us-gaap/role/ref/otherTransitionRef> -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference 6: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 924 -SubTopic 10 -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SAB Topic 11.L) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479941/924-10-599-1Reference 7: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 5 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-5Reference 8: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 30 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-30Reference 9: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 42 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-42Reference 10: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22Reference 11: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 40 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-40Reference 12: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22Reference 13: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-4 + Details Name: us-gaap RevenueFromContractWithCustomerIncludingAssessedTax Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - Details Name: us-gaap ConcentrationRiskByBenchmarkAxis=nuvb CustomerBMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap ConcentrationRiskByBenchmarkAxis=nuvb CustomerBMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 40 R33.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Significant Accounting Policies - Schedule of Concentration of Customers and Suppliers Accounts Receivable (Details) \$ in Thousands Jun. 30, 2024 USD (\$) Concentration Risk [Line Items] A Accounts receivable, net of allowance for credit loss \$ 0 Customer B [Member] A Concentration Risk [Line Items] A Accounts receivable, net of allowance for credit loss \$ 117 X - DefinitionAmount of allowance for credit loss on accounts receivable. + ReferencesReference 1: <http://www.xbrl.org/2009/role/commonPracticeRef> -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(5)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478777/944-210-599-1Reference 2: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SX 210.5-02(4)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-599-1Reference 3: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 326 -SubTopic 20 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479344/326-20-45-1Reference 4: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 310 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481962/310-10-50-4Reference 5: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 326 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 13 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479319/326-20-50-13Reference 6: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 326 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 13 -Subparagraph (f) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479319/326-20-50-13 + Details Name: us-gaap AllowanceForDoubtfulAccountsReceivable Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 954 -SubTopic 310 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Publisher FASB -URI https://asc.fasb.org/1943274/2147478785/954-310-50-2 + Details Name: us-gaap ConcentrationRiskLineItems Namespace Prefix: us-gaap_ Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - Details Name: us-gaap ConcentrationRiskLineItems Namespace Prefix: us-gaap_ Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - Details Name: us-gaap AccountsReceivableMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 41 R34.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Acquisition - Additional Information (Details) \$ in Millions 6 Months Ended Jun. 30, 2024 USD (\$) shares AnHeart [Member] A Business Acquisition [Line Items] A Assets acquired | \$ \$ 450.9 Liabilities assumed | \$ 53.7 Net cash consideration | \$ 425.1 Research and development asset | \$ \$ 425.1 Class A Common Stock [Member] A Business Acquisition [Line Items] A Share issued 27,646,255 Class A Common Stock [Member] | AnHeart [Member] A Business Acquisition [Line Items] A Share issued 27,646,255 Series A Preferred Stock | AnHeart [Member] A Business Acquisition [Line Items] A Share issued 851,202 Convertible Preferred Stock [Member] A Business Acquisition [Line Items] A Share issued 851,202 Convertible Preferred Stock [Member] | AnHeart [Member] A Business Acquisition [Line Items] A Shares issued upon conversion 85,120,200 X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 805 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 5 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479907/805-20-50-5 + Details Name: us-gaap BusinessAcquisitionLineItems Namespace Prefix: us-gaap_ Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - DefinitionAmount of assets acquired at the acquisition date. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Topic 805 -SubTopic 20 -Section 50 -Paragraph 1 -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479907/805-20-50-1 + Details Name: us-gaap BusinessCombinationRecognizedIdentifiableAssetsAcquiredAndLiabilitiesAssumedAssets Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionAmount of liabilities assumed at the acquisition date. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Topic 805 -SubTopic 20 -Section 50 -Paragraph 1 -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479907/805-20-50-1 + Details Name: us-gaap BusinessCombinationRecognizedIdentifiableAssetsAcquiredAndLiabilitiesAssumedLiabilities Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionNumber of shares issued for each share of convertible preferred stock that is converted. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SX 210.5-02(27)) -SubTopic 10 -Topic 210 -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-599-1Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SX 210.5-02(28)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-599-1Reference 3: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Topic 505 -SubTopic 10 -Section 50 -Paragraph 3 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481112/505-10-50-3Reference 4: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 6 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481112/505-10-50-6Reference 5: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 16 -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481112/505-10-50-16 + Details Name: us-gaap ConvertiblePreferredStockSharesIssuedUponConversion Namespace Prefix: us-gaap_ Data Type: xbrli:sharesItem Type Balance Type: na Period Type: instant X - DefinitionThe cash outflow associated with the acquisition of a business, net of the cash acquired from the purchase. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Topic 230 -SubTopic 10 -Section 45 -Paragraph 13 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482740/230-10-45-13 + Details Name: us-gaap PaymentsToAcquireBusinessesNetOfCashAcquired Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionAmount of asset recognized from cost for research and development asset acquired in transaction other than business combination or from joint venture formation or both. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 350 -SubTopic 30 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482665/350-30-50-1 + Details Name: us-gaap ResearchAndDevelopmentAssetAcquiredOtherThanThroughBusinessCombinationFairValueAcquired Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionNumber of new stock issued during the period. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 50 -Paragraph 2 -SubTopic 10 -Topic 505 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481112/505-10-50-2Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SX 210.5-02(28)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-599-1Reference 3: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SX 210.5-02(29)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-599-1Reference 4: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 946 -SubTopic 505 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478448/946-505-50-2Reference 5: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section 599 -Paragraph 3 -Subparagraph (SX 210.6-09(4)(b)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-599-3Reference 6: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 946 -SubTopic 10 -Name Accounting Standards Codification -Section 599 -Paragraph 3 -Subparagraph (SX 210.6-03(i)(1)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479886/946-10-599-3Reference 7: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SX 210.3-04) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480008/505-10-599-1 + Details Name: us-gaap StockIssuedDuringPeriodSharesNewIssues Namespace Prefix: us-gaap_ Data Type: xbrli:sharesItem Type Balance Type: na Period Type: duration X - Details Name: us-gaap BusinessAcquisitionAxis=nuvb AnheartMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap StatementClassOfStockAxis=us-gaap CommonClassAMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap StatementClassOfStockAxis=us-gaap SeriesAPreferredStockMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap StatementClassOfStockAxis=us-gaap ConvertiblePreferredStockMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 42 R35.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Acquisition - Summary of Final Allocable Purchase Price Consists (Details) \$ in Thousands 6 Months Ended Jun. 30, 2024 USD (\$) Business Acquisition [Line Items] A Total allocable purchase price \$ 397,251 Acquisition-related Costs [Member] A Business Acquisition [Line Items] A Total allocable purchase price 7,434 Class A Common Stock [Member] A Business Acquisition [Line Items] A Total allocable purchase price 89,297 [1] Series A Non-Voting Convertible Preferred Stock [Member] A Business Acquisition [Line Items] A Total allocable purchase price 274,938 [2] Consideration Warrants [Member] A Business Acquisition [Line Items] A Total allocable purchase price 764 [3] Assumed Share Based Awards [Member] A Business Acquisition [Line Items] A Total allocable purchase price \$ 24,818 [4] [1] 27,646,255 shares issued at \$3.23 per share as of the Acquisition Date. [2] 851,202 shares issued at as-converted value of \$323 per share. [3] 2,893,731 warrants issued at the estimated fair value of \$0.2641 per warrant. [4] The portion recorded in consideration transferred represents the fair value of replacement awards (Assumed Options & Assumed RSUs) attributable to precombination services. X - DefinitionAllocable Purchase Price + ReferencesNo definition available. + Details Name: nuvb AllocablePurchasePrice Namespace Prefix: nuvb_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 805 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 5 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479907/805-20-50-5 + Details Name: us-gaap BusinessAcquisitionLineItems Namespace Prefix: us-gaap_ Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - Details Name: us-gaap NonrecurringAdjustmentAxis=us-gaap AcquisitionRelatedCostsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap StatementClassOfStockAxis=us-gaap CommonClassAMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details

Name: us-gaap_StatementClassOfStockAxis=us-gaap_ConvertiblePreferredStockMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_StatementClassOfStockAxis=nuvb_ConsiderationWarrantsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_StatementClassOfStockAxis=nuvb_AssumedShareBasedAwardsMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 43 R36.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Acquisition - Schedule Of Final Allocable Purchase Price Consists (Parenthetical) (Details) 6 Months Ended Jun. 30, 2024 \$ / shares shares Business Acquisition [Line Items] A Shares issued, price per share \$ 3.23 Warrants issued [shares 2,893,731 Estimated Fair Value Warrant \$ 0.2641 Class A Common Stock [Member] A Business Acquisition [Line Items] A Share issued [shares 27,646,255 Shares issued, price per share \$ 11.5 Convertible Preferred Stock [Member] A Business Acquisition [Line Items] A Share issued [shares 851,202 Shares issued, price per share \$ 323 X - DefinitionEstimated Fair Value Warrant Per Share + ReferencesNo definition available. + Details Name: nuvb_EstimatedFairValueWarrantPerShare Namespace Prefix: nuvb_ Data Type: dtr-types:perShareItem Type Balance Type: na Period Type: duration X - DefinitionStock and Warrants Issued During Period Share Preferred Stock and Warrants + ReferencesNo definition available. + Details Name: nuvb_StockAndWarrantsIssuedDuringPeriodSharePreferredStockAndWarrants Namespace Prefix: nuvb_ Data Type: xbrli:sharesItem Type Balance Type: na Period Type: duration X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 805 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 5 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479907/805-20-50-5 + Details Name: us-gaap_BusinessAcquisitionLineItems Namespace Prefix: us-gaap_ Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - DefinitionPer share or per unit amount of equity securities issued. + ReferencesNo definition available. + Details Name: us-gaap_SharesIssuedPricePerShare Namespace Prefix: us-gaap_ Data Type: dtr-types:perShareItem Type Balance Type: na Period Type: instant X - DefinitionNumber of new stock issued during the period. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Section 50 -Paragraph 2 -SubTopic 10 -Topic 505 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481112/505-10-50-2Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(28)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 3: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(29)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 505 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/214748448/946-505-50-2Reference 5: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-09(4)(b)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-S99-3Reference 6: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 3 -Subparagraph (SX 210.6-03(i)(1)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479886/946-10-S99-3Reference 7: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.3-04) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480008/505-10-S99-1 + Details Name: us-gaap_StockIssuedDuringPeriodSharesNewIssues Namespace Prefix: us-gaap_ Data Type: xbrli:sharesItem Type Balance Type: na Period Type: duration X - Details Name: us-gaap_StatementClassOfStockAxis=us-gaap_CommonClassAMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_StatementClassOfStockAxis=us-gaap_ConvertiblePreferredStockMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 44 R37.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Fair Value Measurements and Marketable Securities Available For Sale - Schedule of Marketable securities Measured at Fair Value on Recurring Basis (Details) - USD (\$) \$ in Thousands Jun. 30, 2024 Dec. 31, 2023 Cash equivalents: A A Cash equivalents \$ 19,556 \$ 26,509 Marketable securities: A A Marketable securities 542,884 568,564 Total financial assets 562,440 595,073 Financial liabilities: A A Warrants 1,441 353 Money Market Funds [Member] A A Cash equivalents: A A Cash equivalents 7,492 20,574 U.S. Government And Government Agency Securities [Member] A A Cash equivalents: A A Cash equivalents 9,926 4,988 Marketable securities: A A Marketable securities 408,996 453,175 Corporate Bonds [Member] A A Cash equivalents: A A Cash equivalents 2,138 947 Marketable securities: A A Marketable securities 75,639 81,585 Certificate of deposits [Member] A A Marketable securities: A A Marketable securities 21,694 10,176 Commercial Paper [Member] A A Marketable securities: A A Marketable securities 36,555 23,628 Level 1 [Member] A A Cash equivalents: A A Cash equivalents 7,492 20,574 Marketable securities: A A Marketable securities 0 0 Total financial assets 7,492 20,574 Financial liabilities: A A Warrants 1,363 338 Level 1 [Member] | Money Market Funds [Member] A A Cash equivalents: A A Cash equivalents 7,492 20,574 Level 1 [Member] | U.S. Government And Government Agency Securities [Member] A A Cash equivalents: A A Cash equivalents 0 0 Marketable securities: A A Marketable securities 0 0 Level 1 [Member] | Corporate Bonds [Member] A A Cash equivalents: A A Cash equivalents 0 0 Marketable securities: A A Marketable securities 0 0 Level 1 [Member] | Certificate of deposits [Member] A A Marketable securities: A A Marketable securities 0 0 Level 1 [Member] | Commercial Paper [Member] A A Marketable securities: A A Marketable securities 0 0 Level 2 [Member] A A Cash equivalents: A A Cash equivalents 12,064 5,935 Marketable securities: A A Marketable securities 542,884 568,564 Total financial assets 554,948 574,499 Financial liabilities: A A Warrants 0 0 Level 2 [Member] | Money Market Funds [Member] A A Cash equivalents: A A Cash equivalents 0 0 Level 2 [Member] | U.S. Government And Government Agency Securities [Member] A A Cash equivalents: A A Cash equivalents 9,926 4,988 Marketable securities: A A Marketable securities 408,996 453,175 Level 2 [Member] | Corporate Bonds [Member] A A Cash equivalents: A A Cash equivalents 2,138 947 Marketable securities: A A Marketable securities 75,639 81,585 Level 2 [Member] | Certificate of deposits [Member] A A Marketable securities: A A Marketable securities 21,694 10,176 Level 2 [Member] | Commercial Paper [Member] A A Marketable securities: A A Marketable securities 36,555 23,628 Level 3 [Member] A A Cash equivalents: A A Cash equivalents 0 0 Marketable securities: A A Marketable securities 0 0 Total financial assets 0 0 Financial liabilities: A A Warrants 78 15 Level 3 [Member] | Money Market Funds [Member] A A Cash equivalents: A A Cash equivalents 0 0 Level 3 [Member] | U.S. Government And Government Agency Securities [Member] A A Cash equivalents: A A Cash equivalents 0 0 Marketable securities: A A Marketable securities 0 0 Level 3 [Member] | Corporate Bonds [Member] A A Cash equivalents: A A Cash equivalents 0 0 Marketable securities: A A Marketable securities 0 0 Level 3 [Member] | Certificate of deposits [Member] A A Marketable securities: A A Marketable securities 0 0 Level 3 [Member] | Commercial Paper [Member] A A Marketable securities: A A Marketable securities \$ 0 \$ 0 X - DefinitionTotal financial assets: + ReferencesNo definition available. + Details Name: nuvb_TotalFinancialAssets Namespace Prefix: nuvb_ Data Type: xbrli:monetaryItem Type Balance Type: debit Period Type: instant X - ReferencesNo definition available. + Details Name: us-gaap_CashAndCashEquivalentsAbstract Namespace Prefix: us-gaap_ Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - DefinitionAmount of short-term, highly liquid investments that are both readily convertible to known amounts of cash and so near their maturity that they present insignificant risk of changes in value because of changes in interest rates. Excludes cash and cash equivalents within disposal group and discontinued operation. + ReferencesReference 1: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(2)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478777/944-210-S99-1Reference 2: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(1)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1 + Details Name: us-gaap_CashEquivalentsAtCarryingValue Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItem Type Balance Type: debit Period Type: instant X - DefinitionFair value, before effects of master netting arrangements, of a financial liability or contract with one or more underlyings, notional amount or payment provision or both, and the contract can be net settled by means outside the contract or delivery of an asset. Includes liabilities elected not to be offset. Excludes liabilities not subject to a master netting arrangement. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 815 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4B -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480434/815-10-50-4BReference 2: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 815 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4B -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480434/815-10-50-4BReference 3: http://www.xbrl.org/2003/role/exampleRef -Topic 210 -SubTopic 20 -Name Accounting Standards Codification -Section 55 -Paragraph 22 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483444/210-20-55-22Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Topic 210 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483466/210-20-50-3Reference 5: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 210 -SubTopic 20 -Name Accounting Standards Codification -Section 55 -Paragraph 7 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483444/210-20-55-7Reference 6: http://fasb.org/us-gaap/role/ref/legacyRef -Name Accounting Standards Codification -Section 50 -Paragraph 4D -Subparagraph (c)(2) -SubTopic 20 -Topic 860 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481326/860-20-50-4D + Details Name: us-gaap_DerivativeFairValueOfDerivativeLiability Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItem Type Balance Type: credit Period Type: instant X - ReferencesNo definition available. + Details Name: us-gaap_FinancialLiabilitiesFairValueDisclosureAbstract Namespace Prefix: us-gaap_ Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - DefinitionAmount of investment in marketable security. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 942 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.9-03(4)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478546/942-210-S99-1Reference 2: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(2)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1 + Details Name: us-gaap_MarketableSecurities Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItem Type Balance Type: debit Period Type: instant X - ReferencesNo definition available. + Details Name: us-gaap_MarketableSecuritiesAbstract Namespace Prefix: us-gaap_ Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - Details Name: us-gaap_FinancialInstrumentAxis=us-gaap_MoneyMarketFundsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_FinancialInstrumentAxis=us-gaap_CorporateDebtSecuritiesMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_FinancialInstrumentAxis=nuvb_CertificateOfDepositsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_FinancialInstrumentAxis=us-gaap_CommercialPaperMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_FairValueByFairValueHierarchyLevelAxis=us-gaap_FairValueInputsLevel1Member Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_FairValueByFairValueHierarchyLevelAxis=us-gaap_FairValueInputsLevel2Member Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_FairValueByFairValueHierarchyLevelAxis=us-gaap_FairValueInputsLevel3Member Namespace Prefix: Data Type: na Balance Type: Period Type: XML 45 R38.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Fair Value Measurements and Marketable Securities Available For Sale - Schedule of Cost, Aggregate Fair Value and Unrealized Gains of Marketable Securities Available-for-Sale (Details) - USD (\$) \$ in Thousands Jun. 30, 2024 Dec. 31, 2023 Debt Securities, Available-for-Sale, Amortized Cost [Abstract] A A Amortized Cost \$ 563,854 \$ 594,892 Unrealized Gains 64 1,193 Unrealized Losses (1,478) (1,012) Fair Value 562,440 595,073 Cash Equivalents [Member] A A Debt Securities, Available-for-Sale, Amortized Cost [Abstract] A A Amortized Cost 19,556 26,508 Unrealized Gains 0 1 Unrealized Losses 0 0 Fair Value 19,556 26,509 Marketable securities [Member] A A Debt Securities, Available-for-Sale, Amortized Cost [Abstract] A A Amortized Cost 544,298 568,384 Unrealized Gains 64 1,192 Unrealized Losses (1,478) (1,012) Fair Value 542,884 568,564 Money Market Funds [Member] | Cash Equivalents [Member] A A Debt Securities, Available-for-Sale, Amortized Cost [Abstract] A A Amortized Cost 7,492 20,574 Unrealized Gains 0 0 Unrealized Losses 0 0 Fair Value 7,492 20,574 Certificate of deposits [Member] | Marketable securities [Member] A A Debt Securities, Available-for-Sale, Amortized Cost [Abstract] A A Amortized Cost 21,689 10,171 Unrealized Gains 8 7 Unrealized Losses (3) (2) Fair Value 21,694 10,176 Commercial Paper [Member] | Marketable securities [Member] A A Debt Securities, Available-for-Sale, Amortized Cost [Abstract] A A Amortized Cost 36,580 23,609 Unrealized Gains 4 22 Unrealized Losses (29) (3) Fair Value 36,555 23,628 U.S. Government Securities [Member] | Cash Equivalents [Member] A A Debt Securities, Available-for-Sale, Amortized Cost [Abstract] A A Amortized Cost 9,926 4,988 Unrealized Gains 0 0 Unrealized Losses 0 0 Fair Value 9,926 4,988 U.S. Government And Government Agency Securities [Member] | Cash Equivalents [Member] A A Debt Securities, Available-for-Sale, Amortized Cost [Abstract] A A Amortized Cost 4,987 Unrealized Gains 1 1 Unrealized Losses 0 0 Fair Value 4,988 U.S. Government And Government Agency Securities [Member] | Marketable securities [Member] A A Debt Securities, Available-for-Sale, Amortized Cost [Abstract] A A Amortized Cost 410,234 452,813 Unrealized Gains 10 1,095 Unrealized Losses (1,248) (733) Fair Value 408,996 453,175 Corporate Bonds [Member] | Cash Equivalents [Member] A A Debt Securities, Available-for-Sale, Amortized Cost [Abstract] A A Amortized Cost 2,138 947 Unrealized Gains 0 0 Unrealized Losses 0 0 Fair Value 2,138 947 Corporate Bonds [Member] | Marketable securities [Member] A A Debt Securities, Available-for-Sale, Amortized Cost [Abstract] A A Amortized Cost 75,795 81,791 Unrealized Gains 42 68 Unrealized Losses (198) (274) Fair Value \$ 75,639 \$ 81,585 X - DefinitionAmount, before tax, of unrealized gain in accumulated other comprehensive income (AOCI) on investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available-for-sale). + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-2 + Details Name: us-gaap_AvailableForSaleDebtSecuritiesAccumulatedGrossUnrealizedGainBeforeTax Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItem Type Balance Type: credit Period Type: instant X - DefinitionAmount, before tax, of unrealized loss in accumulated other comprehensive income (AOCI) on investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available-for-sale). + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-2 + Details Name: us-gaap_AvailableForSaleDebtSecuritiesAccumulatedGrossUnrealizedLossBeforeTax Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItem Type Balance Type: debit Period Type: instant X - DefinitionAmortized cost of investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available-for-sale). + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(1)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478777/944-210-S99-1Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-2Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 326 -SubTopic 30 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479130/326-30-45-1 + Details Name: us-gaap_AvailableForSaleDebtSecuritiesAmortizedCostBasis Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItem Type Balance Type: debit Period Type: instant X - DefinitionAmount of investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available-for-sale). + ReferencesReference 1: http://www.xbrl.org/2003/role/exampleRef -Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 103 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482078/820-10-55-103Reference 2: http://www.xbrl.org/2003/role/exampleRef -Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 100 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482078/820-10-55-100Reference 3: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (bbb)(2)(i) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2Reference 4: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (bbb)(2) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2Reference 5: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (bbb)(1) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2Reference 6: http://www.xbrl.org/2003/role/disclosureRef -Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2Reference 7: http://www.xbrl.org/2003/role/disclosureRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (aa) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-2Reference 8: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -

Section 45 -Paragraph 1 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481830/320-10-45-1Reference 9: http://www.xbrl.org/2003/role/disclosureRef -Topic 942 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.9-03(6)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478546/942-210-S99-1Reference 10: http://www.xbrl.org/2003/role/disclosureRef -Topic 326 -SubTopic 30 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479130/326-30-45-1 + Details Name: us-gaap AvailableForSaleSecuritiesDebtSecurities Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - ReferencesNo definition available. + Details Name: us-gaap DebtSecuritiesAvailableForSaleAmortizedCostAbstract Namespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - Details Name: us-gaap CashAndCashEquivalentsAxis=us-gaap CashEquivalentsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap CashAndCashEquivalentsAxis=nuvb MarketableSecuritiesMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap FinancialInstrumentAxis=nuvb CertificateOfDepositsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap FinancialInstrumentAxis=us-gaap CommercialPaperMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap FinancialInstrumentAxis=us-gaap USGovernmentDebtSecuritiesMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap FinancialInstrumentAxis=us-gaap USTreasuryAndGovernmentMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap FinancialInstrumentAxis=us-gaap CorporateDebtSecuritiesMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 46 R39.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Fair Value Measurements and Marketable Securities Available for Sale - Schedule of Net Realized Gain (Loss) on Marketable Securities (Details) - USD (\$) \$ in Thousands 3 Months Ended 6 Months Ended Jun. 30, 2024 Jun. 30, 2023 Jun. 30, 2024 Jun. 30, 2023 Net realized gains (losses) on available-for-sale securities were as follows: A A A Realized gains from sales of available-for-sale securities \$ 1 \$ 12 \$ 1 \$ 12 Realized losses from sales of available-for-sale securities (8) (11) (7) (207) Net recognized gains (losses) on equity securities were as follows: A A A Total losses on marketable securities \$ (7) \$ (99) \$ (6) \$ (195) X - DefinitionAmount of realized gain on investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available-for-sale). + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 9 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-9 + Details Name: us-gaap DebtSecuritiesAvailableForSaleRealizedGain Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionAmount of realized loss on investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available-for-sale). + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 9 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-9 + Details Name: us-gaap DebtSecuritiesAvailableForSaleRealizedLoss Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: duration X - ReferencesNo definition available. + Details Name: us-gaap EquitySecuritiesFvNiRealizedGainLossAbstract Namespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - ReferencesNo definition available. + Details Name: us-gaap MarketableSecuritiesGainLossAbstract Namespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionAmount of realized and unrealized gain (loss) on investment in marketable security. + ReferencesNo definition available. + Details Name: us-gaap MarketableSecuritiesGainLossExcludingOtherThanTemporaryImpairments Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration XML 47 R40.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Fair Value Measurements and Marketable Securities Available for Sale - Schedule of Marketable Securities with Continuous Unrealized Losses (Details) - USD (\$) \$ in Thousands Jun. 30, 2024 Dec. 31, 2023 Debt Securities, Held-to-Maturity, Allowance for Credit Loss [Line Items] A A Less than 12 Months, Fair Value \$ 353,295 \$ 236,190 Less than 12 Months, Unrealized Losses (730) (727) 12 Months or Greater, Fair Value 127,265 51,245 12 Months or Greater, Unrealized Losses (748) (284) Debt Securities, Available-for-sale, Unrealized Loss Position 480,560 287,435 Debt Securities, Available-for-sale, Unrealized Loss Position, Accumulated Loss (1,478) (1,011) Certificate of deposits [Member] A A Debt Securities, Held-to-Maturity, Allowance for Credit Loss [Line Items] A A Less than 12 Months, Fair Value 9,940 5,693 Less than 12 Months, Unrealized Losses (3) (2) 12 Months or Greater, Fair Value 0 0 12 Months or Greater, Unrealized Losses 0 0 Debt Securities, Available-for-sale, Unrealized Loss Position 9,940 5,693 Debt Securities, Available-for-sale, Unrealized Loss Position, Accumulated Loss (3) (2) Commercial Paper [Member] A A Debt Securities, Held-to-Maturity, Allowance for Credit Loss [Line Items] A A Less than 12 Months, Fair Value 26,185 9,101 Less than 12 Months, Unrealized Losses (29) (3) 12 Months or Greater, Fair Value 0 0 12 Months or Greater, Unrealized Losses 0 0 Debt Securities, Available-for-sale, Unrealized Loss Position 26,185 9,101 Debt Securities, Available-for-sale, Unrealized Loss Position, Accumulated Loss (29) (3) U.S. Government And Government Agency Securities [Member] A A Debt Securities, Held-to-Maturity, Allowance for Credit Loss [Line Items] A A Less than 12 Months, Fair Value 270,609 199,552 Less than 12 Months, Unrealized Losses (628) (694) 12 Months or Greater, Fair Value 115,683 24,761 12 Months or Greater, Unrealized Losses (620) (38) Debt Securities, Available-for-sale, Unrealized Loss Position 386,292 224,313 Debt Securities, Available-for-sale, Unrealized Loss Position, Accumulated Loss (1,248) (732) Corporate Bonds [Member] A A Debt Securities, Held-to-Maturity, Allowance for Credit Loss [Line Items] A A Less than 12 Months, Fair Value 46,561 21,844 Less than 12 Months, Unrealized Losses (70) (28) 12 Months or Greater, Fair Value 11,582 26,484 12 Months or Greater, Unrealized Losses (128) (246) Debt Securities, Available-for-sale, Unrealized Loss Position 58,143 48,328 Debt Securities, Available-for-sale, Unrealized Loss Position, Accumulated Loss \$ (198) \$ (274) X - DefinitionAmount of investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available-for-sale), in continuous unrealized loss position for more than 12 months, without allowance for credit loss. Includes beneficial interest in securitized financial asset. + ReferencesReference 1: http://www.xbrl.org/2003/role/exampleRef -Topic 326 -SubTopic 30 -Name Accounting Standards Codification -Section 55 -Paragraph 8 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479081/326-30-55-8Reference 2: http://fasb.org/us-gaap/role/ref/otherTransitionRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 7 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-7Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 326 -SubTopic 30 -Name Accounting Standards Codification -Section 50 -Paragraph 5 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479106/326-30-50-5 + Details Name: us-gaap DebtSecuritiesAvailableForSaleContinuousUnrealizedLossPosition12MonthsOrLonger Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionAmount of accumulated unrealized loss on investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available-for-sale), in continuous unrealized loss position for 12 months or longer, without allowance for credit loss. Includes beneficial interest in securitized financial asset. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/otherTransitionRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 7 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-7Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 326 -SubTopic 30 -Name Accounting Standards Codification -Section 50 -Paragraph 5 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479106/326-30-50-5 + Details Name: us-gaap DebtSecuritiesAvailableForSaleContinuousUnrealizedLossPosition12MonthsOrLongerAccumulatedLoss Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionAmount of investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available-for-sale), in continuous unrealized loss position for less than 12 months, without allowance for credit loss. Includes beneficial interest in securitized financial asset. + ReferencesReference 1: http://www.xbrl.org/2003/role/exampleRef -Topic 326 -SubTopic 30 -Name Accounting Standards Codification -Section 55 -Paragraph 8 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479081/326-30-55-8Reference 2: http://fasb.org/us-gaap/role/ref/otherTransitionRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 7 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-7Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 326 -SubTopic 30 -Name Accounting Standards Codification -Section 50 -Paragraph 7 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-7Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 326 -SubTopic 30 -Name Accounting Standards Codification -Section 50 -Paragraph 7 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479106/326-30-50-5 + Details Name: us-gaap DebtSecuritiesAvailableForSaleContinuousUnrealizedLossPositionLessThan12Months Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionAmount of accumulated unrealized loss on investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available-for-sale), in continuous unrealized loss position for less than 12 months, without allowance for credit loss. Includes beneficial interest in securitized financial asset. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/otherTransitionRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 7 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-7Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 326 -SubTopic 30 -Name Accounting Standards Codification -Section 50 -Paragraph 5 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479106/326-30-50-5 + Details Name: us-gaap DebtSecuritiesAvailableForSaleContinuousUnrealizedLossPositionAccumulatedLoss Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesReference 1: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 326 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 13 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479319/326-20-50-13 + Details Name: us-gaap DebtSecuritiesHeldtomaturityAllowanceForCreditLossLineItems Namespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - Details Name: us-gaap FinancialInstrumentAxis=nuvb CertificateOfDepositsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap FinancialInstrumentAxis=us-gaap CommercialPaperMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap FinancialInstrumentAxis=us-gaap CorporateDebtSecuritiesMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 48 R41.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Fair Value Measurements and Marketable Securities Available For Sale - Schedule Of Available for sale Securities Maturity on Fair Value (Details) \$ in Thousands Jun. 30, 2024 USD (\$) Available-for-Sale Securities, Debt Maturities [Abstract] A Due With in one year, Amortized Cost \$ 396,133 Due after one year through five years, Amortized Cost 146,751 Total, Amortized Cost 542,884 Certificate of deposits [Member] A Available-for-Sale Securities, Debt Maturities [Abstract] A Due With in one year, Amortized Cost 21,069 Due after one year through five years, Amortized Cost 625 Total, Amortized Cost 21,694 Commercial Paper [Member] A Available-for-Sale Securities, Debt Maturities [Abstract] A Due With in one year, Amortized Cost 36,555 Due after one year through five years, Amortized Cost 0 Total, Amortized Cost 36,555 U.S. Government And Government Agency Securities [Member] A Available-for-Sale Securities, Debt Maturities [Abstract] A Due With in one year, Amortized Cost 283,225 Due after one year through five years, Amortized Cost 125,771 Total, Amortized Cost 408,996 Corporate Bonds [Member] A Available-for-Sale Securities, Debt Maturities [Abstract] A Due With in one year, Amortized Cost 55,284 Due after one year through five years, Amortized Cost 20,355 Total, Amortized Cost \$ 75,639 X - ReferencesNo definition available. + Details Name: us-gaap AvailableForSaleSecuritiesDebtMaturitiesAbstract Namespace Prefix: us-gaap Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionAmortized cost of investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available-for-sale), with single maturity date and allocated without single maturity date, maturing in second through fifth fiscal year following current fiscal year. Excludes interim and annual periods when interim periods are reported from current statement of financial position date (rolling approach). + ReferencesReference 1: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-3 + Details Name: us-gaap AvailableForSaleSecuritiesDebtMaturitiesAfterOneThroughFiveYearsAmortizedCost Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionAmortized cost of investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available-for-sale), with single maturity date and allocated without single maturity date. + ReferencesReference 1: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-3Reference 2: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-2Reference 3: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 942 -SubTopic 320 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Publisher FASB -URI https://asc.fasb.org/1943274/2147477268/942-320-50-3 + Details Name: us-gaap AvailableForSaleSecuritiesDebtMaturitiesSingleMaturityDateAmortizedCostBasis Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionAmortized cost of investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available-for-sale), with single maturity date and allocated without single maturity date, maturing in next fiscal year following current fiscal year. Excludes interim and annual periods when interim periods are reported from current statement of financial position date (rolling approach). + ReferencesReference 1: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 320 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147481800/320-10-50-3 + Details Name: us-gaap AvailableForSaleSecuritiesDebtMaturitiesWithinOneYearAmortizedCost Namespace Prefix: us-gaap Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - Details Name: us-gaap FinancialInstrumentAxis=nuvb CertificateOfDepositsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap FinancialInstrumentAxis=us-gaap CommercialPaperMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap FinancialInstrumentAxis=us-gaap USTreasuryAndGovernmentMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap FinancialInstrumentAxis=us-gaap CorporateDebtSecuritiesMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 49 R42.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Collaboration and License Agreements - Schedule of revenue recognized (Details) - USD (\$) \$ in Thousands 3 Months Ended 6 Months Ended Jun. 30, 2023 Jun. 30, 2024 Jun. 30, 2023 Revenue \$ 1,435 \$ 0 \$ 1,435 \$ 0 Research and development service revenue A A A Revenue \$ 1,435 A \$ 1,435 A X - DefinitionAmount, excluding tax collected from customer, of revenue from satisfaction of performance obligation by transferring promised good or service to customer. Tax collected from customer is tax assessed by governmental authority that is both imposed on and concurrent with specific revenue-producing transaction, including, but not limited to, sales, use, value added and excise. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 41 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-41Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 270 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -

Paragraph 1 -Subparagraph (i) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482964/270-10-50-1Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (ee) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference 4: http://fasb.org/us-gaap/role/ref/otherTransitionRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference 5: http://fasb.org/us-gaap/role/ref/otherTransitionRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference 6: http://www.xbrl.org/2003/role/disclosureRef -Topic 924 -SubTopic 10 -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SAB Topic 11.L) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479941/924-10-599-1Reference 7: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 5 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-5Reference 8: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 30 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-30Reference 9: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 42 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-42Reference 10: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22Reference 11: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 40 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-40Reference 12: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22Reference 13: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-4 + Details Name: us-gaap RevenueFromContractWithCustomerExcludingAssessedTax Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - Details Name: srt ProductOrServiceAxis=nuvb ResearchAndDevelopmentServiceRevenueMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 50 R43.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Collaboration and License Agreements (Additional Information) (Details) - USD (\$) in Thousands 3 Months Ended 6 Months Ended Jun. 30, 2024 Jun. 30, 2024 Jun. 30, 2023 Jun. 30, 2023 Revenue \$ 1,435 \$ 0 \$ 1,435 \$ 0 Accounts receivable 117 Á \$ 117 Á \$ 0 Revenue, performance obligation satisfied over time, method used, description Á Á The Company expects to recognize all of this balance as revenue over the next 4 years. Á Á Contract Liability 12,200 Á \$ 12,200 Á Á Contract with customer liability revenue recognized 1,300 Á Á Á Á Capitalized contract cost 1,300 Á 1,300 Á Á NK [Member] Á Á Á Á Accounts receivable 100 Á 100 Á Á Research and development service revenue Á Á Á Á Revenue 1,435 Á 1,435 Á Á Contract Liability 21,357 Á 21,357 Á Á Research and development service revenue | NK [Member] Á Á Á Á Revenue 300 Á Á Á Research and development service revenue | Innovent Á Á Á Á Revenue 1,100 Á Á Á Accounts receivable \$ 0 Á \$ 0 Á Á X - DefinitionAmount, after allowance for credit loss, of right to consideration from customer for product sold and service rendered in normal course of business. + ReferencesReference 1: http://www.xbrl.org/2003/role/exampleRef -Topic 946 -SubTopic 310 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Subparagraph (d) -Publisher FASB -URI https://asc.fasb.org/1943274/2147477802/946-310-45-1Reference 2: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(5)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478777/944-210-599-1Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 210 -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SX 210.6-04(5)(b)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479170/946-210-599-1Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Topic 954 -SubTopic 310 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479196/954-310-45-1Reference 5: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(3)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478777/944-210-599-1 + Details Name: us-gaap AccountsReceivableNet Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionAmount, after accumulated amortization and accumulated impairment loss, of asset recognized from cost incurred to obtain or fulfill contract with customer. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 340 -SubTopic 40 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479483/340-40-50-3 + Details Name: us-gaap CapitalizedContractCostNet Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: debit Period Type: instant X - DefinitionAmount of obligation to transfer good or service to customer for which consideration has been received or is receivable. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479837/606-10-45-1Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 8 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-8Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 2 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479837/606-10-45-2 + Details Name: us-gaap ContractWithCustomerLiability Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionAmount of revenue recognized that was previously included in balance of obligation to transfer good or service to customer for which consideration from customer has been received or is due. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 8 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-8 + Details Name: us-gaap ContractWithCustomerLiabilityRevenueRecognized Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionAmount, excluding tax collected from customer, of revenue from satisfaction of performance obligation by transferring promised good or service to customer. Tax collected from customer is tax assessed by governmental authority that is both imposed on and concurrent with specific revenue-producing transaction, including, but not limited to, sales, use, value added and excise. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 41 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-41Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 270 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Subparagraph (i) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482964/270-10-50-1Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (ee) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference 4: http://fasb.org/us-gaap/role/ref/otherTransitionRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference 5: http://fasb.org/us-gaap/role/ref/otherTransitionRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 32 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-32Reference 6: http://www.xbrl.org/2003/role/disclosureRef -Topic 924 -SubTopic 10 -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SAB Topic 11.L) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479941/924-10-599-1Reference 7: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 5 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-5Reference 8: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 30 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-30Reference 9: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 42 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-42Reference 10: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22Reference 11: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 40 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-40Reference 12: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22Reference 13: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-4 + Details Name: us-gaap RevenueFromContractWithCustomerExcludingAssessedTax Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionDescription of method used to recognize revenue for performance obligation that is satisfied over time. Includes, but not is limited to, output or input method. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 18 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-18 + Details Name: us-gaap RevenuePerformanceObligationSatisfiedOverTimeMethodUsedDescription Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - Details Name: us-gaap RelatedPartyTransactionAxis=nuvb NKMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: srt ProductOrServiceAxis=nuvb ResearchAndDevelopmentServiceRevenueMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap RelatedPartyTransactionAxis=nuvb InnoventMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 51 R44.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Collaboration and License Agreements - Schedule Of Contract Assets And Contract Liabilities (Details) \$ in Thousands Jun. 30, 2024 USD (\$) Contract Liability \$ 12,200 Research and development service revenue Á Contract Liability \$ 21,357 X - DefinitionAmount of obligation to transfer good or service to customer for which consideration has been received or is receivable. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 1 -Publisher FASB -URI https://asc.fasb.org/1943274/2147479837/606-10-45-1Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 8 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-8Reference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 42 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-42Reference 4: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22Reference 11: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 40 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-40Reference 12: http://www.xbrl.org/2003/role/disclosureRef -Topic 280 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 22 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482810/280-10-50-22Reference 13: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-4 + Details Name: us-gaap RevenueFromContractWithCustomerExcludingAssessedTax Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionDescription of method used to recognize revenue for performance obligation that is satisfied over time. Includes, but not is limited to, output or input method. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 606 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 18 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479806/606-10-50-18 + Details Name: us-gaap RevenuePerformanceObligationSatisfiedOverTimeMethodUsedDescription Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - Details Name: us-gaap RelatedPartyTransactionAxis=nuvb NKMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: srt ProductOrServiceAxis=nuvb ResearchAndDevelopmentServiceRevenueMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 52 R45.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Borrowings (Additional Information) (Details) - USD (\$) in Millions 3 Months Ended Jun. 30, 2024 Apr. 04, 2023 Loan and Security Agreement [Member] Á Á Subordinated Borrowing [Line Items] Á Á Fixed interest rate on borrowings 4.25% Á Repayments of short term borrowin 0.7 Á Short term bank loan Á \$ 40.0 Short term working capital Á 20.0 Long term debt Á \$ 20.0 Outstanding line of credit balance net \$ 1.8 Á Minimum [Member] | Loan and Security Agreement [Member] Á Á Subordinated Borrowing [Line Items] Á Á Long term debt maturity date Á Apr. 04, 2024 Maximum [Member] | Loan and Security Agreement [Member] Á Á Subordinated Borrowing [Line Items] Á Á Long term debt maturity date Á Apr. 04, 2025 Borrowing In 2020 [Member] | Loan Agreements With Bank of Hangzhou [Member] Á Á Subordinated Borrowing [Line Items] Á Á Fixed interest rate on borrowings 4.05% Á Short-term borrowings outstanding balance \$ 7.0 Á Repayments of short term borrowin 5.6 Á Short term borrowings drawn down \$ 7.0 Á Borrowing In 2022 [Member] | Loan Agreements With China Merchants Bank [Member] Á Á Subordinated Borrowing [Line Items] Á Á Fixed interest rate on borrowings 3.90% Á Short-term borrowings outstanding balance \$ 2.8 Á X - DefinitionLong-Term Borrowin + ReferencesNo definition available. + Details Name: nuvb LongTermBorrowing Namespace Prefix: nuvb_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionShort-Term Borrowings Drawn Down + ReferencesNo definition available. + Details Name: nuvb ShortTermBorrowingsDrawnDown Namespace Prefix: nuvb_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionShort-Term Working Capital + ReferencesNo definition available. + Details Name: nuvb ShortTermWorkingCapital Namespace Prefix: nuvb_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionAmount of short-term bank loan secured by broker-dealer customer's security. + ReferencesReference 1: http://www.xbrl.org/2009/role/commonPracticeRef -Name Rule 15c3-1 -Number 240 -Section 15c3-1 -Publisher SEC + Details Name: srt BankLoans Namespace Prefix: srt_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionMaturity date of long-term debt, in YYYY-MM-DD format. + ReferencesReference 1: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (bbb)(2)(i) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2Reference 2: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (bbb)(2) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2Reference 3: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (bbb)(1) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2 + Details Name: us-gaap LongTermDebtMaturityDate Namespace Prefix: us-gaap_ Data Type: xbrli:dateTimeType Balance Type: na Period Type: instant X - DefinitionAmount of cash outflow for payment of an obligation from a lender, including but not limited to, letter of credit, standby letter of credit and revolving credit arrangements. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 230 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 15 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482740/230-10-45-15Reference 2: http://www.xbrl.org/2009/role/commonPracticeRef -Topic 235 -SubTopic 10 -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SX 210.4-08(f)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480678/235-10-599-1 + Details Name: us-gaap RepaymentsOfLinesOfCredit Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionThe cash outflow for a borrowing having initial term of repayment within one year or the normal operating cycle, if longer. + ReferencesReference 1: http://fasb.org/us-gaap/role/ref/legacyRef -Topic 230 -SubTopic 10 -Name Accounting Standards Codification -Section 45 -Paragraph 15 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482740/230-10-45-15 + Details Name: us-gaap RepaymentsOfShortTermDebt Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionFor the form of debt having an initial term of less than one year or less than the normal operating cycle, if longer, average borrowings during the period. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Name Regulation S-K (SK) -Number 229 -Section 1402 -Paragraph a -Publisher SECReference 2: http://www.xbrl.org/2003/role/disclosureRef -Name Regulation S-K (SK) -Number 229 -Section 1402 -Paragraph b -Subparagraph (1) -Publisher SECReference 3: http://www.xbrl.org/2003/role/disclosureRef -Topic 946 -SubTopic 220 -Name Accounting Standards Codification -Section 599 -Paragraph 1 -Subparagraph (SX 210.6-07(3)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147479134/946-220-599-1 + Details Name: us-gaap ShorttermDebtAverageOutstandingAmount Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionStated interest rate of the subordinated debt. + ReferencesNo definition available. + Details Name: us-gaap SubordinatedBorrowingInterestRate Namespace Prefix: us-gaap_ Data Type: dtr-types:percentItemType Balance Type: na Period Type: duration X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesNo definition available. + Details Name: us-gaap SubordinatedBorrowingLineItems Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - Details Name: us-gaap TypeOfArrangementAxis=nuvb LoanAndSecurityAgreementMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: srt RangeAxis=srt_MinimumMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap SubordinatedBorrowingAxis=nuvb BorrowingTwoThousandsAndTwentyMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap TypeOfArrangementAxis=nuvb LoanAgreementsWithBankOfHangzhouMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap SubordinatedBorrowingAxis=nuvb BorrowingTwoThousandsAndTwentyTwoMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap TypeOfArrangementAxis=nuvb LoanAgreementsWithChinaMerchantsBankMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 53 R46.htm IDEA: XBRL

DOCUMENT v3.24.2.u1 Leases - Schedule of Aggregate Lease Payments (Details) \$ in Thousands Jun. 30, 2024 USD (\$) Leases [Abstract] Å 2024 (remaining 6 months) \$ 1,095 2025 1,416 2026 629 2027 and thereafter 81 Total undiscounted lease payments 3,221 Less: imputed interest (173) Total operating lease liabilities \$ 3,048 X - ReferencesNo definition available. + Details Name: us-gaap LeasesAbstract Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionAmount of lessee's undiscounted obligation for lease payment for operating lease. + ReferencesReference 1:

[Member] | Maximum [Member] | Class A Shares [Member] A Share Based Compensation Arrangement By Share Based Payment Award [Line Items] A Number of common shares issuable under the plan (in shares) 4,750,354 X - Definition Number of shares issuable under share based payment arrangement. + ReferencesNo definition available. + Details Name: nuvb NumberOfSharesIssuableUnderShareBasedPaymentArrangement Namespace Prefix: nuvb Data Type: xbrli:sharesItem Type Balance Type: na Period Type: duration X - DefinitionAggregate number of common shares reserved for future issuance. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(29)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1> + Details Name: us-gaap CommonStockCapitalSharesReservedForFutureIssuance Namespace Prefix: us-gaap Data Type: xbrli:sharesItem Type Balance Type: na Period Type: instant X - DefinitionDescription of terms of share-based payment arrangement. Includes, but is not limited to, type of award or grantee and reason for issuance. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Topic 718 -SubTopic 10 -Section 50 -Paragraph 1 -Subparagraph (a) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-1> -Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Topic 718 -SubTopic 10 -Section 50 -Paragraph 2 -Subparagraph (a) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> + Details Name: us-gaap ShareBasedCompensationArrangementByShareBasedPaymentAwardDescription Namespace Prefix: us-gaap Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 35 -Paragraph 1D -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480483/718-10-35-1> -Reference 2: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 35 -Paragraph 3 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480483/718-10-35-3> -Reference 3: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (a)(1) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 4: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (a)(2) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 5: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (a)(3) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 6: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (a)(1)(i) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 7: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(ii) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 8: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(iii) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 9: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(iv) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 10: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(iv)(01) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 11: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(iv)(02) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 12: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(iv)(03) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 13: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(iv)(04) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 14: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(2)(i) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 15: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(2)(ii) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 16: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(2)(iii) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 17: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(2)(iii)(01) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 18: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(2)(iii)(02) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 19: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(2)(iii)(03) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 20: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (d)(1) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 21: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (d)(2) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 22: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (e)(1) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 23: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (e)(2) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 24: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (f)(2)(i) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 25: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (f)(2)(ii) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 26: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (f)(2)(iii) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 27: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (f)(2)(iv) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 28: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (f)(2)(v) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> + Details Name: us-gaap ShareBasedCompensationArrangementByShareBasedPaymentAwardLineItems Namespace Prefix: us-gaap Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - DefinitionWeighted average grant-date fair value of options vested. + ReferencesNo definition available. + Details Name: us-gaap ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsVestedWeightedAverageGrantDateFairValue Namespace Prefix: us-gaap Data Type: dtr-types:perShareItem Type Balance Type: na Period Type: duration X - DefinitionPurchase price of common stock expressed as a percentage of its fair value. + ReferencesNo definition available. + Details Name: us-gaap SharebasedCompensationArrangementBySharebasedPaymentAwardPurchasePriceOfCommonStockPercent Namespace Prefix: us-gaap Data Type: dtr-types:percentItem Type Balance Type: na Period Type: duration X - Details Name: us-gaap PlanNameAxis=nuvb TwoThousandAndTwentyOneEquityIncentivePlanMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: srt RangeAxis=srt MaximumMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap StatementClassOfStockAxis=us-gaap CommonClassAMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap PlanNameAxis=nuvb EmployeeStockPurchasePlanMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 58 R51.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Stock-Based Compensation - Summary of Stock-Based Compensation Expense (Details) - USD (\$) in Thousands 3 Months Ended 6 Months Ended Jun 30, 2024 Jun 30, 2023 Jun 30, 2021 Jun 30, 2023 Employee Service Share Based Compensation Allocation Of Recognized Period Costs [Line Items] A A A Stock-based compensation expense \$ 9,569 \$ 5,523 \$ 14,610 \$ 10,360 Research and Development [Member] A A A Employee Service Share Based Compensation Allocation Of Recognized Period Costs [Line Items] A A A Stock-based compensation expense 4,190 2,420 6,482 4,431 General and Administrative [Member] A A A Employee Service Share Based Compensation Allocation Of Recognized Period Costs [Line Items] A A A Stock-based compensation expense \$ 5,379 \$ 3,103 \$ 8,128 \$ 5,929 X - DefinitionAmount of expense for award under share-based payment arrangement. Excludes amount capitalized. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SAB Topic 14.F) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147479830/718-10-S99-1> -Reference 2: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (h)(1)(i) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> + Details Name: us-gaap AllocatedShareBasedCompensationExpense Namespace Prefix: us-gaap Data Type: xbrli:monetaryItem Type Balance Type: debit Period Type: duration X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesNo definition available. + Details Name: us-gaap EmployeeServiceShareBasedCompensationAllocationOfRecognizedPeriodCostsLineItems Namespace Prefix: us-gaap Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - Details Name: us-gaap IncomeStatementLocationAxis=us-gaap ResearchAndDevelopmentExpenseMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap IncomeStatementLocationAxis=us-gaap SellingGeneralAndAdministrativeExpensesMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 59 R52.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Stock-Based Compensation - Summary of Stock Option Activity under 2019 Plan and 2021 plan (Details) - 2021 and 2019 Equity Incentive Plan [Member] - shares 6 Months Ended Jun 30, 2024 Share-Based Payment Arrangement, Expensed and Capitalized, Amount [Line Items] A Outstanding at December 31, 2023 30,649,239 Options assumed for the Acquisition of AnHeart 13,742,239 Granted 11,322,679 Exercised (1,798,190) Forfeited (1,223,019) Outstanding at June 30, 2024 52,692,948 Exercisable at June 30, 2024 23,909,107 X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesNo definition available. + Details Name: us-gaap EmployeeServiceShareBasedCompensationAllocationOfRecognizedPeriodCostsLineItems Namespace Prefix: us-gaap Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - DefinitionThe number of shares into which fully or partially vested stock options outstanding as of the balance sheet date can be currently converted under the option plan. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(iii) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> + Details Name: us-gaap ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsExercisableNumber Namespace Prefix: us-gaap Data Type: xbrli:sharesItem Type Balance Type: na Period Type: instant X - DefinitionThe number of shares under options that were cancelled during the reporting period as a result of occurrence of a terminating event specified in contractual agreements pertaining to the stock option plan. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(iv)(03) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> + Details Name: us-gaap ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsForfeituresInPeriod Namespace Prefix: us-gaap Data Type: xbrli:sharesItem Type Balance Type: na Period Type: duration X - DefinitionGross number of share options (or share units) granted during the period. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(iv)(01) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> + Details Name: us-gaap ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsGrantsInPeriodGross Namespace Prefix: us-gaap Data Type: xbrli:sharesItem Type Balance Type: na Period Type: duration X - DefinitionNumber of options outstanding, including both vested and non-vested options. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(i) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 2: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(ii) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> + Details Name: us-gaap ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsOutstandingNumber Namespace Prefix: us-gaap Data Type: xbrli:sharesItem Type Balance Type: na Period Type: instant X - DefinitionNumber of shares of stock issued during the period pursuant to acquisitions. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(28)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1> -Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(29)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1> -Reference 3: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.3-04) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480008/505-10-S99-1> + Details Name: us-gaap StockIssuedDuringPeriodSharesAcquisitions Namespace Prefix: us-gaap Data Type: xbrli:sharesItem Type Balance Type: na Period Type: duration X - DefinitionNumber of share options (or share units) exercised during the current period. + ReferencesReference 1: <http://fasb.org/us-gaap/role/ref/legacyRef> -Name Accounting Standards Codification -Section 50 -Paragraph 2 -SubTopic 10 -Topic 505 -Publisher FASB -URI <https://asc.fasb.org/1943274/2147481112/505-10-50-2> -Reference 2: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(28)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1> -Reference 3: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(29)) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480566/210-10-S99-1> -Reference 4: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(iv)(02) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> -Reference 5: <http://fasb.org/us-gaap/role/ref/legacyRef> -Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.3-04) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480008/505-10-S99-1> + Details Name: us-gaap StockIssuedDuringPeriodSharesStockOptionsExercised Namespace Prefix: us-gaap Data Type: xbrli:sharesItem Type Balance Type: na Period Type: duration X - Details Name: us-gaap PlanNameAxis=nuvb TwoThousandAndTwentyOneEquityIncentivePlanAndTwoThousandAndNineteenEquityIncentivePlanMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 60 R53.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Stock-Based Compensation - Summary of Restricted Stock Unit Activity (Details) - Restricted Stock Unit [Member] 6 Months Ended Jun 30, 2024 shares Share-Based Payment Arrangement, Expensed and Capitalized, Amount [Line Items] A Non-vested at December 31, 2023 0 RSUs assumed for the Acquisition of AnHeart 2,201,694 Granted 0 Vested (40,342) Forfeited or canceled 0 Non-vested at June 30, 2024 2,161,352 X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesNo definition available. + Details Name: us-gaap EmployeeServiceShareBasedCompensationAllocationOfRecognizedPeriodCostsLineItems Namespace Prefix: us-gaap Data Type: xbrli:stringItem Type Balance Type: na Period Type: duration X - DefinitionGross number of share options (or share units) granted during the period. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef> -Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (c)(1)(iv)(01) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480429/718-10-50-2> + Details Name: us-gaap ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsGrantsInPeriodGross Namespace Prefix: us-gaap Data Type: xbrli:sharesItem Type Balance Type: na Period Type: duration X - DefinitionNumber of non-vested options outstanding. + ReferencesNo definition available. + Details Name: us-gaap SharebasedCompensationArrangementBySharebasedPaymentAwardOptionsNonvestedNumberOfShares Namespace Prefix: us-gaap Data Type: xbrli:sharesItem Type Balance Type: na Period Type: instant X - DefinitionNumber of non-vested options forfeited. + ReferencesNo definition available. + Details Name: us-

gaap_SharebasedCompensationArrangementBySharebasedPaymentAwardOptionsNonvestedOptionsForfeitedNumberOfShares Namespace Prefix: us-gaap_ Data Type: xbrli:sharesItemType Balance Type: na Period Type: duration X - DefinitionNumber of options vested. + ReferencesNo definition available. + Details Name: us-gaap_SharebasedCompensationArrangementBySharebasedPaymentAwardOptionsVestedNumberOfShares Namespace Prefix: us-gaap_ Data Type: xbrli:sharesItemType Balance Type: na Period Type: duration X - DefinitionNumber of shares of stock issued during the period pursuant to acquisitions. + ReferencesReference 1: [http://fasb.org/us-gaap/role/ref/legacyRef-Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.5-02\(28\)\)](http://fasb.org/us-gaap/role/ref/legacyRef-Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(28))) -Publisher FASB -URI [https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef-Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.5-02\(29\)\)](https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef-Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(29))) -Publisher FASB -URI [https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 3: http://fasb.org/us-gaap/role/ref/legacyRef-Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.5-02\(29\)\)](https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 3: http://fasb.org/us-gaap/role/ref/legacyRef-Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(29))) -Publisher FASB -URI <https://asc.fasb.org/1943274/2147480008/505-10-S99-1> + Details Name: us-gaap_StockIssuedDuringPeriodSharesAcquisitions Namespace Prefix: us-gaap_ Data Type: xbrli:sharesItemType Balance Type: na Period Type: duration X - Details Name: us-gaap_AwardTypeAxis=us-gaap_RestrictedStockUnitsRSUMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 61 R54.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Warrants -Additional Information (Details) \$ / shares in Units, \$ in Millions 3 Months Ended 6 Months Ended Jun. 30, 2024 USD (\$) \$ / shares shares Jun. 30, 2024 \$ / shares shares Class Of Warrant Or Right [Line Items] A Shares issued, price per share | \$ / shares \$ 3.23 \$ 3.23 Change in Fair Value, Gain (Loss) | \$ \$ 1.1 A Fair Value Equity Level1 To Level2 or Level3 Transfers Description A no Public Warrants [Member] A Class Of Warrant Or Right [Line Items] A Warrants to purchase common stock outstanding 4,791,639 4,791,639 Class of Warrant or Right, Number of Securities Called by Warrants or Rights 4,791,639 4,791,639 Private Placement Warrants [Member] A Class Of Warrant Or Right [Line Items] A Warrants to purchase common stock outstanding 162,500 162,500 Class of Warrant or Right, Number of Securities Called by Warrants or Rights 162,500 162,500 Forward Purchase Warrants [Member] A Class Of Warrant Or Right [Line Items] A Warrants to purchase common stock outstanding 833,333 833,333 Class of Warrant or Right, Number of Securities Called by Warrants or Rights 833,333 833,333 Class A Common Stock [Member] A Class Of Warrant Or Right [Line Items] A Warrants to purchase common stock outstanding 5,787,472 5,787,472 Aggregate warrants outstanding 8,681,193 8,681,193 Shares issued, price per share | \$ / shares \$ 11.5 \$ 11.5 Class of Warrant or Right, Number of Securities Called by Warrants or Rights 5,787,472 5,787,472 Class A Common Stock [Member] | Consideration Warrants [Member] A Class Of Warrant Or Right [Line Items] A Warrants to purchase common stock outstanding 2,893,731 2,893,731 Class of Warrant or Right, Number of Securities Called by Warrants or Rights 2,893,731 2,893,731 X - DefinitionFair Value Equity Level1 To Level2 or Level3 Transfers Description + ReferencesNo definition available. + Details Name: nuvb_FairValueEquityLevel1ToLevel2OrLevel3TransfersDescription Namespace Prefix: nuvb_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesNo definition available. + Details Name: us-gaap_ClassOfWarrantOrRightLineItems Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionNumber of securities into which the class of warrant or right may be converted. For example, but not limited to, 500,000 warrants may be converted into 1,000,000 shares. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef-Topic 505 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481112/505-10-50-3> + Details Name: us-gaap_ClassOfWarrantOrRightNumberOfSecuritiesCalledByWarrantsOrRights Namespace Prefix: us-gaap_ Data Type: xbrli:sharesItemType Balance Type: na Period Type: instant X - DefinitionNumber of warrants or rights outstanding. + ReferencesNo definition available. + Details Name: us-gaap_ClassOfWarrantOrRightOutstanding Namespace Prefix: us-gaap_ Data Type: xbrli:sharesItemType Balance Type: na Period Type: instant X - DefinitionAmount of gain (loss) from (increase) decrease in fair value of contract or contract feature in long-duration contract issued by insurance entity that both protects contract holder from other-than-nominal capital market risk and exposes insurance entity to other-than-nominal capital market risk. Excludes change attributable to instrument-specific credit risk. + ReferencesReference 1: <http://www.xbrl.org/2003/role/disclosureRef-Topic 944 -SubTopic 40 -Name Accounting Standards Codification -Section 45 -Paragraph 3 -Publisher FASB -URI https://asc.fasb.org/1943274/2147480112/944-40-45-3> + Details Name: us-gaap_MarketRiskBenefitChangeInFairValueGainLoss Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionPer share or per unit amount of equity securities issued. + ReferencesNo definition available. + Details Name: us-gaap_SharesIssuedPricePerShare Namespace Prefix: us-gaap_ Data Type: dtr-types:perShareItemType Balance Type: na Period Type: instant X - Details Name: us-gaap_ClassOfWarrantOrRightAxis=nuvb_PublicWarrantsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_ClassOfWarrantOrRightAxis=nuvb_PrivatePlacementWarrantsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_ClassOfWarrantOrRightAxis=nuvb_ForwardPurchaseWarrantsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_StatementClassOfStockAxis=us-gaap_CommonClassAMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_ClassOfWarrantOrRightAxis=nuvb_ConsiderationWarrantsMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 62 R55.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Warrants -Schedule of Quantitative Information Regarding Level 3 Fair Value Measurements (Details) -Level 3 [Member] - \$ / shares 6 Months Ended 12 Months Ended Jun. 30, 2024 Dec. 31, 2023 Class of Warrant or Right [Line Items] A Share Price \$ 2.92 \$ 1.51 Strike price \$ 11.5 \$ 11.5 Expected term in years 1 year 7 months 6 days 2 years 1 month 6 days Expected volatility 98.10% 82.70% Risk-free interest rate 4.90% 4.10% Dividend 0.00% 0.00% X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesNo definition available. + Details Name: us-gaap_ClassOfWarrantOrRightLineItems Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionPer share amount received by subsidiary or equity investee for each share of common stock issued or sold in the stock transaction. + ReferencesNo definition available. + Details Name: us-gaap_SaleOfStockPricePerShare Namespace Prefix: us-gaap_ Data Type: dtr-types:perShareItemType Balance Type: na Period Type: instant X - DefinitionThe estimated dividend rate (a percentage of the share price) to be paid (expected dividends) to holders of the underlying shares over the option's term. + ReferencesReference 1: [http://www.xbrl.org/2003/role/disclosureRef-Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph \(f\)\(2\)\(iii\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2](http://www.xbrl.org/2003/role/disclosureRef-Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (f)(2)(iii) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2) + Details Name: us-gaap_ShareBasedCompensationArrangementByShareBasedPaymentAwardFairValueAssumptionsExpectedDividendRate Namespace Prefix: us-gaap_ Data Type: dtr-types:percentItemType Balance Type: na Period Type: duration X - DefinitionThe estimated measure of the percentage by which a share price is expected to fluctuate during a period. Volatility also may be defined as a probability-weighted measure of the dispersion of returns about the mean. The volatility of a share price is the standard deviation of the continuously compounded rates of return on the share over a specified period. That is the same as the standard deviation of the differences in the natural logarithms of the stock prices plus dividends, if any, over the period. + ReferencesReference 1: [http://www.xbrl.org/2003/role/disclosureRef-Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph \(f\)\(2\)\(ii\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2](http://www.xbrl.org/2003/role/disclosureRef-Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (f)(2)(ii) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2) + Details Name: us-gaap_ShareBasedCompensationArrangementByShareBasedPaymentAwardFairValueAssumptionsExpectedVolatilityRate Namespace Prefix: us-gaap_ Data Type: dtr-types:percentItemType Balance Type: na Period Type: duration X - DefinitionThe risk-free interest rate assumption that is used in valuing an option on its own shares. + ReferencesReference 1: [http://www.xbrl.org/2003/role/disclosureRef-Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph \(f\)\(2\)\(iv\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2](http://www.xbrl.org/2003/role/disclosureRef-Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (f)(2)(iv) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2) + Details Name: us-gaap_ShareBasedCompensationArrangementByShareBasedPaymentAwardFairValueAssumptionsRiskFreeInterestRate Namespace Prefix: us-gaap_ Data Type: dtr-types:percentItemType Balance Type: na Period Type: duration X - DefinitionType of a single share of a number of saleable stocks of a company. + ReferencesNo definition available. + Details Name: us-gaap_SharePrice Namespace Prefix: us-gaap_ Data Type: dtr-types:perShareItemType Balance Type: na Period Type: instant X - DefinitionExpected term of award under share-based payment arrangement, in 'PnYmMnDTnHnMnS' format, for example, 'P1Y5M13D' represents reported fact of one year, five months, and thirteen days. + ReferencesReference 1: [http://www.xbrl.org/2003/role/disclosureRef-Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph \(f\)\(2\)\(i\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2](http://www.xbrl.org/2003/role/disclosureRef-Topic 718 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (f)(2)(i) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480429/718-10-50-2) + Details Name: us-gaap_ShareBasedCompensationArrangementBySharebasedPaymentAwardFairValueAssumptionsExpectedTerm1 Namespace Prefix: us-gaap_ Data Type: xbrli:durationItemType Balance Type: na Period Type: duration X - Details Name: us-gaap_FairValueByFairValueHierarchyLevelAxis=us-gaap_FairValueInputsLevel3Member Namespace Prefix: Data Type: na Balance Type: Period Type: XML 63 R56.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Warrants -Schedule of Fair Values for Outstanding Warrants (Details) \$ in Thousands Jun. 30, 2024 USD (\$) Class Of Warrant Or Right [Line Items] A Fair values of outstanding warrants \$ 1,441 Public Warrants [Member] A Class Of Warrant Or Right [Line Items] A Fair values of outstanding warrants 767 Private Placement Warrants [Member] A Class Of Warrant Or Right [Line Items] A Fair values of outstanding warrants 78 Forward Purchase Warrants [Member] A Class Of Warrant Or Right [Line Items] A Fair values of outstanding warrants 133 Consideration Warrants [Member] A Class Of Warrant Or Right [Line Items] A Fair values of outstanding warrants \$ 463 X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesNo definition available. + Details Name: us-gaap_ClassOfWarrantOrRightLineItems Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionValue of outstanding derivative securities that permit the holder the right to purchase securities (usually equity) from the issuer at a specified price. + ReferencesReference 1: [http://www.xbrl.org/2009/role/commonPracticeRef-Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph \(bbb\)\(2\) \(i\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2Reference 2: http://www.xbrl.org/2009/role/commonPracticeRef-Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph \(bbb\)\(2\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2Reference 3: http://www.xbrl.org/2009/role/commonPracticeRef-Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph \(bbb\)\(1\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2](http://www.xbrl.org/2009/role/commonPracticeRef-Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (bbb)(2) (i) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2Reference 2: http://www.xbrl.org/2009/role/commonPracticeRef-Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (bbb)(2) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2Reference 3: http://www.xbrl.org/2009/role/commonPracticeRef-Topic 820 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Subparagraph (bbb)(1) -Publisher FASB -URI https://asc.fasb.org/1943274/2147482106/820-10-50-2) + Details Name: us-gaap_WarrantsAndRightsOutstanding Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - Details Name: us-gaap_ClassOfWarrantOrRightAxis=nuvb_PublicWarrantsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_ClassOfWarrantOrRightAxis=nuvb_PrivatePlacementWarrantsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_ClassOfWarrantOrRightAxis=nuvb_ForwardPurchaseWarrantsMember Namespace Prefix: Data Type: na Balance Type: Period Type: X - Details Name: us-gaap_ClassOfWarrantOrRightAxis=nuvb_ConsiderationWarrantsMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 64 R57.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Warrants -Summary of Changes in Liabilities Classified in Level 3 of the Fair Value Hierarchy (Details) \$ in Thousands 6 Months Ended Jun. 30, 2024 USD (\$) Warrants and Rights Note Disclosure [Abstract] A Beginning balance \$ 15 Change in fair value of Private Warrants liability recognized in earnings 63 Ending balance \$ 78 X - DefinitionChange in fair value of private warrants liability recognized in earnings. + ReferencesNo definition available. + Details Name: nuvb_ChangeInFairValueOfPrivateWarrantsLiabilityRecognizedInEarnings Namespace Prefix: nuvb_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: duration X - DefinitionChanges in liabilities classified in Level 3 of fair value hierarchy. + ReferencesNo definition available. + Details Name: nuvb_ChangesInLiabilitiesClassifiedInLevel3OfFairValueHierarchy Namespace Prefix: nuvb_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - ReferencesNo definition available. + Details Name: us-gaap_WarrantsAndRightsNoteDisclosureAbstract Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration XML 65 R58.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Commitments and Contingencies -Additional Information (Details) -Standby Letter of Credit [Member] \$ in Millions 6 Months Ended Jun. 30, 2024 USD (\$) Loss Contingencies [Line Items] A Letter of Credit, Amount \$ 0.6 Line of credit facility, frequency of commitment fee payment annually X - DefinitionThe carrying value as of the balance sheet date of the current and noncurrent portions of long-term obligations drawn from a line of credit, which is a bank's commitment to make loans up to a specific amount. Examples of items that might be included in the application of this element may consist of letters of credit, standby letters of credit, and revolving credit arrangements, under which borrowings can be made up to a maximum amount as of any point in time conditional on satisfaction of specified terms before, as of and after the date of drawdowns on the line. Includes short-term obligations that would normally be classified as current liabilities but for which (a) postbalance sheet date issuance of a long term obligation to refinance the short term obligation on a long term basis, or (b) the enterprise has entered into a financing agreement that clearly permits the enterprise to refinance the short-term obligation on a long term basis and the following conditions are met (1) the agreement does not expire within 1 year and is not cancelable by the lender except for violation of an objectively determinable provision, (2) no violation exists at the BS date, and (3) the lender has entered into the financing agreement is expected to be financially capable of honoring the agreement. + ReferencesReference 1: [http://fasb.org/us-gaap/role/ref/legacyRef-Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.5-02\(22\)\) -SubTopic 10 -Topic 210 -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef-Topic 942 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.9-03\(16\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478546/942-210-S99-1Reference 3: http://www.xbrl.org/2009/role/commonPracticeRef-Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.7-03\(a\)\(16\)\(a\)\(2\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478777/944-210-S99-1](http://fasb.org/us-gaap/role/ref/legacyRef-Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(22)) -SubTopic 10 -Topic 210 -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef-Topic 942 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.9-03(16)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478546/942-210-S99-1Reference 3: http://www.xbrl.org/2009/role/commonPracticeRef-Topic 944 -SubTopic 210 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.7-03(a)(16)(a)(2)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147478777/944-210-S99-1) + Details Name: us-gaap_LineOfCreditNamespaces Namespace Prefix: us-gaap_ Data Type: xbrli:monetaryItemType Balance Type: credit Period Type: instant X - DefinitionDescription of when the fee for available but unused credit capacity is required to be paid (for example, annually, quarterly, or monthly). + ReferencesReference 1: [http://fasb.org/us-gaap/role/ref/legacyRef-Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.5-02\(22\)\(b\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef-Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph \(SX 210.5-02\(19\)\(b\)\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1](http://fasb.org/us-gaap/role/ref/legacyRef-Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(22)(b)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1Reference 2: http://fasb.org/us-gaap/role/ref/legacyRef-Topic 210 -SubTopic 10 -Name Accounting Standards Codification -Section S99 -Paragraph 1 -Subparagraph (SX 210.5-02(19)(b)) -Publisher FASB -URI https://asc.fasb.org/1943274/2147480566/210-10-S99-1) + Details Name: us-gaap_LineOfCreditFacilityFrequencyOfCommitmentFeePayment Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - DefinitionLine items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table. + ReferencesReference 1: [http://www.xbrl.org/2009/role/commonPracticeRef-Topic 450 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483076/450-20-50-4Reference 2: http://www.xbrl.org/2009/role/commonPracticeRef-Topic 450 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 9 -Subparagraph \(a\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483076/450-20-50-9Reference 3: http://www.xbrl.org/2003/role/disclosureRef-Topic 720 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483359/720-20-50-1Reference 4: http://www.xbrl.org/2003/role/exampleRef-Topic 460 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 27 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482395/460-10-55-27Reference 5: http://www.xbrl.org/2003/role/disclosureRef-Topic 450 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 9 -Subparagraph \(b\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483076/450-20-50-9Reference 6: http://www.xbrl.org/2003/role/disclosureRef-Topic 450 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph \(b\) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483076/450-20-50-4Reference 7: http://www.xbrl.org/2003/role/disclosureRef-Topic 460 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482425/460-10-50-3Reference 8: http://www.xbrl.org/2003/role/disclosureRef-Topic 450 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483076/450-20-50-1](http://www.xbrl.org/2009/role/commonPracticeRef-Topic 450 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483076/450-20-50-4Reference 2: http://www.xbrl.org/2009/role/commonPracticeRef-Topic 450 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 9 -Subparagraph (a) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483076/450-20-50-9Reference 3: http://www.xbrl.org/2003/role/disclosureRef-Topic 720 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483359/720-20-50-1Reference 4: http://www.xbrl.org/2003/role/exampleRef-Topic 460 -SubTopic 10 -Name Accounting Standards Codification -Section 55 -Paragraph 27 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482395/460-10-55-27Reference 5: http://www.xbrl.org/2003/role/disclosureRef-Topic 450 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 9 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483076/450-20-50-9Reference 6: http://www.xbrl.org/2003/role/disclosureRef-Topic 450 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 4 -Subparagraph (b) -Publisher FASB -URI https://asc.fasb.org/1943274/2147483076/450-20-50-4Reference 7: http://www.xbrl.org/2003/role/disclosureRef-Topic 460 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 3 -Publisher FASB -URI https://asc.fasb.org/1943274/2147482425/460-10-50-3Reference 8: http://www.xbrl.org/2003/role/disclosureRef-Topic 450 -SubTopic 20 -Name Accounting Standards Codification -Section 50 -Paragraph 1 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483076/450-20-50-1) + Details Name: us-gaap_LossContingenciesLineItems Namespace Prefix: us-gaap_ Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - Details Name: us-gaap_LossContingenciesByNatureOfContingencyAxis=us-gaap_StandbyLettersOfCreditMember Namespace Prefix: Data Type: na Balance Type: Period Type: XML 66 R59.htm IDEA: XBRL DOCUMENT v3.24.2.u1 Subsequent Event (Additional

Information) (Details) Jun. 30, 2024 shares Common Class A [Member] A SubsequentEvent [Line Items] A Warrant exercisable 8,681,193 X - Definition Number of warrants or rights outstanding. + References No definition available. + Details Name: us-gaap_ClassOfWarrantOrRightOutstanding Namespace Prefix: us-gaap_Data Type: xbrli:sharesItemType Balance Type: na Period Type: instant X - DefinitionDetail information of subsequent event by type. User is expected to use existing line items from elsewhere in the taxonomy as the primary line items for this disclosure, which is further associated with dimension and member elements pertaining to a subsequent event. + ReferencesReference 1: http://www.xbrl.org/2003/role/disclosureRef -Topic 830 -SubTopic 30 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Publisher FASB -URI https://asc.fasb.org/1943274/2147481674/830-30-50-2Reference 2: http://www.xbrl.org/2003/role/disclosureRef -Topic 855 -SubTopic 10 -Name Accounting Standards Codification -Section 50 -Paragraph 2 -Publisher FASB -URI https://asc.fasb.org/1943274/2147483399/855-10-50-2 + Details Name: us-gaap_SubsequentEventLineItems Namespace Prefix: us-gaap_Data Type: xbrli:stringItemType Balance Type: na Period Type: duration X - Details Name: us-gaap_StatementClassOfStockAxis=us-gaap_CommonClassAMember Namespace Prefix: Data Type: na Balance Type: Period Type: EXCEL 67 Financial_Report.xlsx IDEA: XBRL DOCUMENT begin 644 Financial_Report.xlsx M4\$#L#110 ((%%5\$D'04UBO +\$ 0 9&Jc4')O<!,O87P+GAM M&SV/_OL/!,1\$ _!QO:+=!P4)B0+12L+ (>/>QLO8D1#1LD) O<CGLCVX>;QA&WPIG M*N*1#BV&5 C(II/>147BK9 7;=N')=H16-Y *OGDK7A.YNJQ<+&4GPZAA!OW J=0U[R;UEA 6#M17E!+ P04 M =>11059'9'>O>I K @&# 1Q08U[R;W1\$+V-O&UL&GVP/ M.L.P#=(?9B>R/F\$S44R:2M.D+PQ6V-CV&IK&B?&UDCJZJ.F-5L#K!'C19 M'0U(2R&DU>'SQ(OT/!1L=9+L+L'..76!CSD!&101!0R9YH'1Q'Q-9'>U'&H00'&#. *E. MH.10 /3ZJ1V7R&GW,K L>JLV7&80 M K'Q5:IMX=-=M%J02P,5% @. @44669E&UL&1\$!>IHXX%[OK]J9 9M'8BV:>'\$W-I=MNTF83M M3A<=&\$5B:~EDD81 OTV23J; / 0LZ?0.14? GZ&AY!<XN8NB&3R M>+>J06N1J+ MUES&6QHOQJ!JM-O=5H11;'&\$81V1?>5XL\$!&505%;U!>M.42, @5RU2-9;.! M&U+1J&KFM(+Y; 7VMX<9<_1.ATR@6XP&U@&? YOI<^1.6HCA5,+&\$P&IG/U9K MQJ'22(''R7V4!;I)J/3%0@R#3ZLC5C.=GSVQ.V7C.K=#1M&N#C17@XML02.MBW A(5M>5 TR M6!'VULS2 Y9>?> IUE!K9;O=05SP6.YXB1'~QL4\$08Q&Z98T91G=4 D. #?# MT4Q0?P!JMHK(P1+27)6SRF9U4H(FLB!J4>'(<7K;YH) M5Z282=J\$~'!&N&M<<~>9ST6S[I25T+7950-RCEU@5 9<8WS2J]-25%UGB5P&M MG#P=P+&+E L&L08:7'82J3E'>34@3 BNEVOXKR2.FJW'52M'/F(9 M-AIRMI1:IMG&IA&L!>L1>[2M!M'6:PUDSY@R.S-D77.U!\$.\$9)->-T'8LZ2 M&D&L'~H8X2IKMHG%8!V/>7L-P>B'N;JN'Z&U3-L+([W1]072N0J)<Z3T MIZ..60FJA%9JGZJ'>ZH<C'Q!?'&Y'CJE>@<@&'Y;A;O&N'@!J;J;~<'K<^+& M.7N'><^<[!GT;7'2MSAD6R4)RU3393>=& M!Y'&V14 5*E=?> EKIDHN'YJ.O.FOHH?0+> Y% GM,T+T.W,F&J# M-M2D& #>?>QK&B=DSN7V85VWGV-1^~?4;[C!SR!6>@!(J[AJ]&&F. # M0X=Y>U>'89Y7&4&04;6RL'Q&M&W'JL \$%&.1DB'V@!X.O40+R4E5!&5O& RN0 MHQO.C37H<~>77% CT9+CVZ9EM6Z0?<9;2)2.<[JF!>GK-J@>?'51W455OR ML+Y>J053L 6:~W6P[13A8+&SDACEL'>FJ2+S&8;~YRMQ'Q&X04T5MA7&R+SC MYL=Q3E.X&G.V#P(RN;LYJ7IE,6>FMM#M L6X9A9N3J7>W5YNTNB42QJ[P# 4A M#W+&C[~^3.G>U _HL@6V\$5#)DU1?>0XGJ/3-R0JA4)?>.NVB8+A=OB5,V[&KXF M8\$O#>FZ=>+?2 VU!4/074;SHYG/JC' MYA.L0Z1~P7V'BH 1Jv~<NJI/>26<.[1N8@F S6VZ3VW>?> 2K6J5DQ\$ M2P=V(Y&8XQ;J1'7CQ1BK;KQ<~>.QOY@CS#%F'~'19H;J6+K&P'F!J M40#EJ046C#30.9FVJ/D3H[W/O#;#Q[!A8N 5!+ P04 M' 11059'9'DG=> &#N)@& 'AL+W=OJ.KCZ)&ET!54CE'~>SV'H&Q1.SEH+&9K9JN]W\$6XB;M QOYV%#&M;YJ,XAHNU;R; M7>?A8454VW.(N<Q X-3K[MVHT:~E=1C&XD:1) (TBKIX0J'CT9WR.VMRX#>8+&YT1Z+=+A=3H;N1<=4L5/X&A\$G 0RDJ; MSEKQ/VXYBY!([EO&5@G6Y<^JL?(<Y;Y.N;Y/6HXID0B&IXT&AS&W,19A:~2@ M+5HJWR-TW@JN>->~E5F'LP\!\$2.9? @J/7BK#5L5\$ >_!&J6[<Q02+&B/ MDV&2 27K ~E<K16>~R'H\$A!%\$0Y Y4U16P\$69Y<T<~>P+8J'PZZQ?(<L#< MC.8ERVQJZJ)/3I5<~>V6>I<7S;N+I&K& M\$>Q'>J7WC8<+&Y9W M~N31VW%YLE~JF!7I7H<621S1)<FJGJY#;9P!>D M\$UMSC&PFJF8&L:D!6JAB2-~>9IHQ<~>[W5L1E&6] MFD:/2Z/AW7J[R#Q>>C\$N!5?D'FY>4Q>NM20!H%\$U 0U+?V#~ -T[64B3JW]T M&= ~E+&2F4=W(E)(TOJ/DJ+GR0~U8W%YWB'DE!851:4J;?=>H9UT+~\$25 M0U1L1=<[!<N4J9VJ63=9EOZ1<~>M'DV~>74FFK45Q+JJOZA'!A=2U6K\$-1F-B: M!F\$ M!F\$ 682)LD X(F)(ITHXKJ)]'FDH#B5%?>M?>D&SF>?0@E-4<+@=(M+>V~N'W! M&@R';V@%&.IOGE)O/F/&OJ. 8A..20>(A49RQ1&*@Q8Z=L3\$<^X=)JN0GZ MH17~T/(YXXYX D6L?G P[P<9) 6QB6~'8S'#6!UOJ]>M'@&B+J>X?<~>QJ]T)=!(-FZY'-L;U.=2 MTS2 (4%[CGT4~M M (DBC/<~>[IJCJ>D>KTR9(B56D MQ'T>~>VTS)0W2JZ'V+VYVC:DWOKJ,F2(E5!O.0J72Z/U,-!#3W%R9S+> MHT@>OQUB83;2JQB8;SS4>90?U]M@&S.GJUNCO&HNCXK M0F\$(A)>QJZV2S+>~.XU2>N!N.MGCSUB*QBQB&B&105W/L&J3H)O=>?>B.KN69W4877L\$&K& M\$>R&F\$N170H? 7D('B)8 M[8H+&ERBX7M\$C&U&L\$PPYBQ'<E"0MBJG'9W'2: M!ZW AJW. 1R8Q&L4'9>HJ)=<D&#F/<~>L'Y?>(V?M>XLO!1%#L+U5H 4S;M M~L1ZYV1?RDJ/RNF:R@>AT71Y-QR<~>S'X/U)/!\$&8@& J]L.F.X0L&4A M#M ,AE#6P;01'446~70!Y~Y6 M.F%(E5V3PU2W)04QRV3.FM3~>('NW(BL/YZ [Q,T,I)ZCQ!E'P\$7 MF\$S1+P'FU01N15)N3C~>WJ/J57'YYQ[QJ]MRF:R M&= ~E+&2F4=W(E)(TOJ/DJ+GR0~U8W%YWB'DE!851:4J;?=>H9UT+~\$25 M0U1L1=<[!<N4J9VJ63=9EOZ1<~>M'DV~>74FFK45Q+JJOZA'!A=2U6K\$-1F-B: M!F\$ M!F\$ 682)LD X(F)(ITHXKJ)]'FDH#B5%?>M?>D&SF>?0@E-4<+@=(M+>V~N'W! 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M&@R';V@%&.IOGE)O/F/&OJ. 8A..20>(A49RQ1&*@Q8Z=L3\$<^X=)JN0GZ MH17~T/(YXXYX D6L?G P[P<9) 6QB6~'8S'#6!UOJ]>M'@&B+J>X?<~>QJ]T)=!(-FZY'-L;U.=2 MTS2 (4%[CGT4~M M (DBC/<~>[IJCJ>D>KTR9(B56D MQ'T>~>VTS)0W2JZ'V+VYVC:DWOKJ,F2(E5!O.0J72Z/U,-!#3W%R9S+> MHT@>OQUB83;2JQB8;SS4>90?U]M@&S.GJUNCO&HNCXK M0F\$(A)>QJZV2S+>~.XU2>N!N.MGCSUB*QBQB&B&105W/L&J3H)O=>?>B.KN69W4877L\$&K& M\$>R&F\$N170H? 7D('B)8 M[8H+&ERBX7M\$C&U&L\$PPYBQ'<E"0MBJG'9W'2: M!ZW AJW. 1R8Q&L4'9>HJ)=<D&#F/<~>L'Y?>(V?M>XLO!1%#L+U5H 4S;M M~L1ZYV1?RDJ/RNF:R@>AT71Y-QR<~>S'X/U)/!\$&8@& J]L.F.X0L&4A M#M ,AE#6P;01'446~70!Y~Y6 M.F%(E5V3PU2W)04QRV3.FM3~>('NW(BL/YZ [Q,T,I)ZCQ!E'P\$7 MF\$S1+P'FU01N15)N3C~>WJ/J57'YYQ[QJ]MRF:R M&= ~E+&2F4=W(E)(TOJ/DJ+GR0~U8W%YWB'DE!851:4J;?=>H9UT+~\$25 M0U1L1=<[!<N4J9VJ63=9EOZ1<~>M'DV~>74FFK45Q+JJOZA'!A=2U6K\$-1F-B: M!F\$ M!F\$ 682)LD X(F)(ITHXKJ)]'FDH#B5%?>M?>D&SF>?0@E-4<+@=(M+>V~N'W! M&@R';V@%&.IOGE)O/F/&OJ. 8A..20>(A49RQ1&*@Q8Z=L3\$<^X=)JN0GZ MH17~T/(YXXYX D6L?G P[P<9) 6QB6~'8S'#6!UOJ]>M'@&B+J>X?<~>QJ]T)=!(-FZY'-L;U.=2 MTS2 (4%[CGT4~M M (DBC/<~>[IJCJ>D>KTR9(B56D MQ'T>~>VTS)0W2JZ'V+VYVC:DWOKJ,F2(E5!O.0J72Z/U,-!#3W%R9S+> MHT@>OQUB83;2JQB8;SS4>90?U]M@&S.GJUNCO&HNCXK M0F\$(A)>QJZV2S+>~.XU2>N!N.MGCSUB*QBQB&B&105W/L&J3H)O=>?>B.KN69W4877L\$&K& M\$>R&F\$N170H? 7D('B)8 M[8H+&ERBX7M\$C&U&L\$PPYBQ'<E"0MBJG'9W'2: M!ZW AJW. 1R8Q&L4'9>HJ)=<D&#F/<~>L'Y?>(V?M>XLO!1%#L+U5H 4S;M M~L1ZYV1?RDJ/RNF:R@>AT71Y-QR<~>S'X/U)/!\$&8@& J]L.F.X0L&4A M#M ,AE#6P;01'446~70!Y~Y6 M.F%(E5V3PU2W)04QRV3.FM3~>('NW(BL/YZ [Q,T,I)ZCQ!E'P\$7 MF\$S1+P'FU01N15)N3C~>WJ/J57'YYQ[QJ]MRF:R M&= ~E+&2F4=W(E)(TOJ/DJ+GR0~U8W%YWB'DE!851:4J;?=>H9UT+~\$25 M0U1L1=<[!<N4J9VJ63=9EOZ1<~>M'DV~>74FFK45Q+JJOZA'!A=2U6K\$-1F-B: M!F\$ M!F\$ 682)LD X(F)(ITHXKJ)]'FDH#B5%?>M?>D&SF>?0@E-4<+@=(M+>V~N'W! 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UJ28&E5#QWUUNW JJK6819QXQT[R]L7F9T6LP7C6A@8?@:K F59VA.SGQ[Q]N8@HLT=07L.[K5QIX6>5 M8Y[KYR9&A:U91 MB-A#PET:~X2B2PFR9.SF =]R[=1SPCA _INS\$%4^)+^" M3FII[AE.H@]~>8B] MN1 KA42A94K9JKB%P5A[F+T]G+V60R.EP:AJ:SGU@/IHE6S_VYMK4V1Y MS).F).47.1\$1C.G.D7K.12^>60.J/J@9T+V7]/.F)XJ(XF V5QYQ4RY19X1?< D=CN2 \$%715).J]. S7J=&49CRU MDZ0 M9K5S64K@556Q^&Z2ID60A:8%9IBC3!5"[.=QPIS: B4QFWS-!\$K M@S@CMC&T2 Z4@366I4U @X(F6~HC9^PD Q%(U MR; Q08<=0%PNB7J15:B.YFQ@>HAEPF@G* PVD)=3.2]7PLSHIMM4W+O+8 MH@ 4PMYY5[S=S2RTH:N#^52HTL7Z)1];7E0HKWYX54@;S.EWFH61=J.A M7?Q&F[BA:5E 71\$~. *7)VT.=W@% U IP9 T9T\$VZV[K]~ U.LK<@69[>97 M]~T/HY[;OCC>WUJ T F[NQ:M1+EY@~^)^K~K[MJ](MOCM0&=O(C):[M=+Z7D]M[~XO3=;@#9J1C3HEO@]/@80H/F5UJY1[OT7T57P02P.% M@ @.44%6?MH@16S^ O84 I.D IX;]JW;K&ULM3W9DMMD+Y^<D3/LI%?>D=-F6%?>D3.2%].%5RS.[L:](% D M88\$ QOZ17 JYED^#G;+; M.@F4V9J[K~RLZLQ@/O:IOUDM91TR=+5=N7)JNN MWV [J]~MV:V766S^7L[F:7].2.%A@]XG6G.C.BAF\$<Y3 @A:~XSTBH^%&T^ !>VPI&X<=D>6H^~ MI^ V2H2)=A*(O19YFA~!8^G@NZ1 ?Z@B!BU05?UIFE)[+&C&SR] M M<2^KJCF@=0?>P.~.IC1COTGZ^ AM5J](S9EEU60>D?I!~.N6-C432~R3C^! M4W@.RKVHFH!T@RHSR5J@~@HJA54.7 VY^OQ4]T]]@JBLAZ~<7I QZ@B E 9CFM MG-ZB^D.190.1^J.KD<@@UN&S^E0^VMN<H@HJ[C]^S^S].N M0 MBID9UWD@>1E^H^~*SR^QC 74JWRS151IBR8T]6%& @JHJA54.7 D510E1+Q5 M7ZQOOTSQ= R4H=AKMYP1[Q]5O1=606%N:XEA819SQSDT\$+DS6M@A^ V4 M3;S7O^V: /IPI;R90>6@I@. M 86I[3=Z3=]C 0J07~RYL~W8@=OV7 M^S^OIJYCPM? 6SDZVIT^75J.7BFOETG1569XJ.7KE<+> *0&7GMEW5C817 M=1S:0I0T~RCP.KWIMF^20A@[OT]^&K& 3UCG 8-KDX<[P@=3@C^TVY^N@ M\$S^<O^O.D~%& (P\$1\$^@/5P-MK=D- &F\$>1-\$!87>352 R3^<@V91.8@ M6EJOF1A.CK.G0&G@T=V@.8= SAJ]/.M\$M+^T14B17@]18%>2LGRK; M=R@I@]YMA. +(#8&2^2^A^L7#++M@B>^S^DRBFO7UA]#9U P\$CQ15YTB^ M@B\$S:~.0XC^M^T+R^W.2 (\$L3^<^>~>~?7IB]T7AB@6A1>3=++^Y^T^M=C^PYD4^.(T4B L^M)^P^J^B^GMEH^Y.XSNFX@^E(B&^S^R0597F^APK ML^ NS>G^D^H^N5;62OVT^M^?42BB^<^I^@07R 3A^E/6;+1H~(M&N~;P>7 M67D%~.LG@BKR28T]2DP.ETHS]JY^P A W59PWK=JW.X2N@M M+1+9&@<@<M^ M6^3TZY0@^MRX1.1\$%&O^F^C3P^>=OASUAY4-H9\$P;~ =#4]@6%P~ 80Q M+1=80>@>A^R]P.U/N2:5\$R@.4^9S^Q307>=+3+B5+55;7L\$Y@I@20IHX^R] MU7<V194I1MM DG@<@.1A(D^+ 71WPD2Y#1 .2F9URG5@E\$S.BRGW=+O8-D@V0 M M+21^HMQJ\$S=ZN;E5R@.4^9@AYW@>+L+=49(T^R^M^RD@G@H^R M9JBJ (3Q.^AR^P2<74^KK@<A1^?~.8=^>1)7 .3^VX.ZX.~4^P@&I53 GML<@88E:~ Q+JX&C^+A]E^@F=O~NH&C^/8@S^M^F5^V^I2.PHLV ZC M1+M MP3]8F@<EGG>+2]^"G [LXKF^+D3.]I79ZUZ<7.2^I~93SE@<^R4UK= MM~^K4+>YR DC5(HJWBV2>~J.X)F/P?> 90DY&XJ5E^1^P?M^L7.KX5I M,5(4#7)U [J]~(Z^HLV ZC M01-4YH1@>V1<D1H55X+W=//7VZ&H163.3^W8)VP@!!;QJ[NM=^]T[C]. MDU^M^YV9^FOS41>=+A1(9RH?)~14^U^I^V+J1AS[R&H;B:6X^D^F5ET]JQ M\$^PYG9Y^Q^<U^<L^U^>M~N~HFDX;V@~B^BZPSZ]3>@Q^TQIROX0 B]@4D^P^V. MLVFS 98Z7UU^ZY5\$VZF+ZR3R PZ8 C6&1.C[?F0R31]^I.C. /F:H@Z1.8Z< MBJ3>2^>[4#&9@J1.VRES<C>3J00+4^=EDCPEF.IQO1Q9[AI]TGKGV5Y0AQ MY^3>PX&G649K8V\$^MNNIR(L0=^@)XWA#< <J^P^U^K:5P[1X]>@&@%<@.OJZ MS.L]I+&C.AXSZ.AQ^O\$[S]JYA~0H48" I"-7M3L8(4UKA)%1HJX\$2@3V M73=5V=P/6.NH=)P]"N^Q6(64-KIR4) %J]S2XPWHU/XS4%-L+^"1&./Z> M 7Y5X 4E16A;J@>?&O^&#B.DS.]ZU^J^N^D="GAODG1I~VSD^LQ PZL3 MC9H8@<2>3^S.Y3N^Y7QOW~F8L.XWCD@E-MG<~2.3^MC M10^D^8RT3(KX@02~)Y^%?JBI^04U!\$O MC=Y(C? +@Y@BCOR...+Z.]<42+Z&R&]JZV Z 8-ME.3T.5VT;PEI]I[1#EYV M8 OSI-GU@<2>3^/SCRS58T^_#K<I@PGK13FC%>?^# .FHE.M;B.P.E+K.C.M 8&K;QYAJ.OA(LI)-> =,LD>7U~W/9+4DZ;P=K\$N(QVY<+<@< M]IJP.@6TU!^ Q^DW\$B\$2AHL& (CHALN~WSLRWA^O^F^F.V.DX6I^M T-O^V M V M^T^~XK3\$^\$V(HI3M3Q051>5>G6JB)S^L^M:~)BS^< <SNX8~Y]LJ@Y48J>M+6P\$SRTP[+X^<=<I&M^F@>+<O^I+W&ASBBJW5&D+ILCY2T4BN^% %XGCMW: M=#\$#KXWQX05EV-D^P^ML^EONCOK@J8B]25Q0J[ENY9PQ6A2]Q5T\$# (MYM3 JG+Y1I&G^H0TJ&6D MKJ..73+=+03E084Q>?>M@?>3J. VF 8MBCVL MOCGT8J)!@=./IMU(V QM?Q5GZ2KL^J./Tl;S^X^V^V:VINOXVH+?SR!M^M/QC!GWYRE3\$4? L]5.ZKTN31Z7CZY04;^<=(+>+>O1@G0^Z^YQ<Z<7^% M11+>+>J^P]J. @4N]J!>+>A#H5LQW353JGJ^?PBY9]!X^EBE^F^=" I]QI]71BY6AQ M^TQ10.4^AN00 M^IR<+&D7 HIL>Y^GS;R[29Q=G 7A3PGP6M]5^Q^R6HWM!5]@FE!IBU3OFL>E1 MY<=&83B.BOKJT4^I~M(3E/2Y.N2LL5A>X^KG&^\$!3%1RDB(\$:W!@Y79X M#>GJXS0+^; +BAJ)4JFY0X41\$5X]SVJ9A7<+&X^>+&J^P^F&A.~>49PQG M3/3LG^<@>^?^ZXSPP0IG6I\$^.(J4KJYI74 M7R1290@46A5 4+>1JOT@ P5574~V^EC74].7@KE AWZU(O>DIT@EY M.W.WT.1JFYLQ&W19 FXEP&SA^N&X(L^MA L^FZ M];.I!)E!Z>N~W%\$[F1C10G<@<@;S.WW.16^CNUY1 Y^Q^Q E6SE^1XX\$5%5.5LTUWWDJ\$~DWHWTM W^P M^T.7J.VVFXB2PM41QRH\$0>@J]I]V O \$A46&XN;5^I.=.T2&[T^T^ M=2U6B+SEM INQMA^6^>+>+>A>+<+>+YSMS>MBCH5A^~IF2IY.~@~63^3Y% M+&C;VVP/@C70.%+&L^H!^A? 7U]7J.GYF@X^39.34?Q@<^Y^X TBFU/D@GFIWU.]80220W.1N^K~.Q-M4<+>=B5&SS8J6BAJCP^W9~L9A\$+X^F7Y;G^#&+^!+>0.59PZD]RLHES M^FAOCI^<L^V].GMM.\$QD QJ/[TQ01K88R.H1D.S@O&F9P<@<5R/8K^&8 MK2+>V. M~LSJQNQ%555YJ]i6&J[P]P2]B]WRNN^AC@BKWQO>90FC MF12G30^QF.#1I59QDQJ5HUKI8IEC~!R!R+&\$WL3I* HYO(].C\$V\$X(8 MPV)](R7<I)CYMV^E+<F(2+42)OM5AEB0+Y^?SSRKH-8>2^H9^KQ. M@Q (CF^R0&+&V9G1S.SUP7^K0.N^JUS8H7DZ043 \$5=V^6VA;D. 3TIT.^KJ04^RAZZ M^CH

[illegible]

[illegible]

[illegible]

YYL8B&/%>_9C@6{/T}>+W.3^*CF^LSEQ1^<+94ZX MT7D)YCA^>+<{UR^Y6G\$HA)RC.0.LY7_R5<f{K NR:|XK63L&D|J4V>S^M^N\$Y&CF>{D&L&C 317RN0.,DN:X58HZU3U-
 8/UXHUYEDYD)3(G\$^X>+>6 M(V?@0(SDC-UQI=7L4S(L:<2?E)ZW^OYT^<2VJ3.E@3I^#0KOLE+68A:0!#L M^0C^@{/R\$S>RE)=S\$6@H^H\$V:W5S0(-
 U49KJH95R9^*Z^M4QZGHD0A|B^A^M\$TRTXG.\$(G@&5\$18CA^*TJ7^>ZE(EM/L^E7(24J0IG|,\$=-_BOX+G4 M>+>3059K,Z+MQ27%>4 1!+|EV0ETO(0>\$\$(\$Y+>.#
 /Z1+>JQ.KL@NJJ^*K MVIFC>|E^I^VS^##K0#fPX.GFWHK7^M.Y<J@05NN^Z^UJG

3CP^3WB6/2W6UI.S4"#GB*4;K'>1HCR%,,/'XL4)-(9;7N+&7*!CR6%-M*,^5LAU043\$!8PZ55GP0,6C*C@8!6!E=&

and Exchange Commission data and sources stated that government employees within the scope of such employment are not subject to domestic corruption prosecution.

U.S.C. 105: var Show={} ; Show.LastAR=null; Show.showAR=function(a,r,w){if(Show.LastAR)Show.hideAR();var e=a;while(e&&e.nodeName!=""TABLE"e=e.nextSibling;if(e.nodeName=="TABLE"e=ref=(window)?w.document.document.getElementById(r):if(ref)=ref.cloneNode(0);e.removeAttribute('id');a.parentNode.appendChild(e)});if(e.style.display=="block"Show.LastAR=e;Show.hideAR=function() {Show.LastAR.style.display="none"};Show.toggleNext=function(a){var e=a;while(e.nodeName!="DIV"e=e.nextSibling;if(e.style){if(e.style.display){if(e.style.display=="none"e=d="block;p="-"}else{d="none;p="+"}e.style.display=d;if(a.textContent){a.textContent=p+a.textContent.substring(1)}else{a.innerHTMLText=p+a.innerHTMLText.substring(1)}}} XML 69 report.css IDEA: XBRL DOCUMENT /* Updated 2009-11-04 */ /* v2.2.0.24 */ /* DefRef Styles */ report table.authRefData { background-color: #def; border: 2px solid #2F4497; font-size: 1em; position: absolute; } report table.authRefData a { display: block; font-weight: bold; } report table.authRefData p { margin-top: 0px; } report table.authRefData .hide { background-color: #2F4497; padding: 1px 3px 0px 0px; text-align: right; } report table.authRefData .hide a: hover { background-color: #2F4497; } report table.authRefData .body { height: 150px; overflow: auto; width: 400px; } report table.authRefData table { font-size: 1em; } /* Report Styles */ .pl.a .pl.a-visited { color: black; text-decoration: none; } /* table */ report { background-color: white; border: 2px solid #acf; clear: both; color: black; font: normal 8pt Helvetica, Arial, sans-serif; margin-bottom: 2em; } report hr { border: 1px solid #acf; } /* Top labels */ report th { background-color: #acf; color: black; font-weight: bold; text-align: center; } report th,td { background-color: transparent; color: #000000; font: bold 10pt Helvetica, Arial, sans-serif; text-align: left; } report .pl { text-align: left; vertical-align: top; white-space: normal; width: 200px; } report .pl { word-wrap: break-word; } report td,pl,a { cursor: pointer; display: block; width: 200px; overflow: hidden; } report td,pl,div,a { width: 200px; } report td,pl,a: hover { background-color: #ffc; } /* Header rows... */ report tr,th { background-color: #acf; color: black; font-weight: bold; } /* Calendars... */ report .rc { background-color: #f0f0f0; } /* Even rows... */ report .re, report .reu { background-color: #def; } report .reu td { border-bottom: 1px solid black; } /* Odd rows... */ report .ro, report .rou { background-color: white; } report .rou td { border-bottom: 1px solid black; } report .rou table td, report .rou table td { border-bottom: 0px solid black; } /* styles for footnote marker */ report .fn { white-space: nowrap; } /* styles for numeric types */ report .num, report .nump { text-align: right; white-space: nowrap; } report .nump { padding-left: 2em; } report .nump { padding: 0px 0.4em 0px 2em; } /* styles for text types */ report .text { text-align: left; white-space: normal; } report .big { margin-bottom: 1em; width: 17em; } report .text .more { display: none; } report .text .note { font-style: italic; font-weight: bold; } report .text .small { width: 10em; } report .sup { font-style: italic; } report .outerFootnotes { font-size: 1em; } XML 71 FilingSummary.xml IDEA: XBRL DOCUMENT 3.24.2.u1.html2292971true670false6 falsefalseR1.htm100000 - Document - Document and Entity InformationSheethttp://nuvationbio.com/20240630/taxonomy/role/DocumentDocumentAndEntityInformationDocument and Entity InformationCover1 falsefalseR2.htm100010 - Statement - Consolidated Balance Sheets (Unaudited)Sheethttp://nuvationbio.com/20240630/taxonomy/role/StatementConsolidatedBalanceSheetsUnauditedConsolidated Balance Sheets (Unaudited)Statements2 falsefalseR3.htm100030 - Statement - Consolidated Balance Sheets (Parenthetical) (Unaudited)Sheethttp://nuvationbio.com/20240630/taxonomy/role/StatementConsolidatedBalanceSheetsParentheticalUnauditedConsolidated Balance Sheets (Parenthetical) (Unaudited)Statements3 falsefalseR4.htm100060 - Statement - Consolidated Statements of Operations and Comprehensive Loss (Unaudited)Sheethttp://nuvationbio.com/20240630/taxonomy/role/StatementConsolidatedStatementsOfOperationsAndComprehensiveLossUnauditedConsolidated Statements of Operations and Comprehensive Loss (Unaudited)Statements4 falsefalseR5.htm100070 - Statement - Consolidated Statements of Redeemable Convertible Preferred Stock and Stockholders' Equity (Unaudited)Sheethttp://nuvationbio.com/20240630/taxonomy/role/StatementConsolidatedStatementsOfRedeemableConvertiblePreferredStockAndStockholdersEquityUnaudited Statements of Redeemable Convertible Preferred Stock and Stockholders' Equity (Unaudited)Statements5 falsefalseR6.htm100080 - Statement - Consolidated Statements of Cash Flows (Unaudited)Sheethttp://nuvationbio.com/20240630/taxonomy/role/StatementConsolidatedStatementsOfCashFlowsUnauditedConsolidated Statements of Cash Flows (Unaudited)Statements6 falsefalseR7.htm995410 - Disclosure - Pay vs Performance DisclosureSheethttp://xbrl.sec.gov/ecd/role/PvpDisclosurePay vs Performance DisclosureNotes7 falsefalseR8.htm995445 - Disclosure - Insider Trading ArrangementsSheethttp://xbrl.sec.gov/ecd/role/InsiderTradingArrangementsInsider Trading ArrangementsNotes8 falsefalseR9.htm995455 - Disclosure - Nature of OperationsSheethttp://nuvationbio.com/20240630/taxonomy/role/Role DisclosureNatureOfOperationsNature of OperationsNotes9 falsefalseR10.htm995465 - Disclosure - Significant Accounting PoliciesSheethttp://nuvationbio.com/20240630/taxonomy/role/Role DisclosureSignificantAccountingPoliciesSignificant Accounting PoliciesNotes10 falsefalseR11.htm995475 - Disclosure - AcquisitionSheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureAcquisitionAcquisitionNotes11 falsefalseR12.htm995485 - Disclosure - Fair Value Measurements and Marketable Securities Available for SaleSheethttp://nuvationbio.com/20240630/taxonomy/role/Role DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSaleFair Value Measurements and Marketable Securities Available for SaleNotes12 falsefalseR13.htm995495 - Disclosure - Collaboration and License AgreementsSheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureCollaborationAndLicenseAgreementsCollaboration and License AgreementsNotes13 falsefalseR14.htm995505 - Disclosure - BorrowingsSheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureBorrowingsBorrowingsNotes14 falsefalseR15.htm995525 - Disclosure - LeasesSheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureLeasesLeasesNotes15 falsefalseR16.htm995535 - Disclosure - Net Loss Per ShareSheethttp://nuvationbio.com/20240630/taxonomy/role/Role DisclosureNetLossPerShareNet Loss Per ShareNotes16 falsefalseR17.htm995545 - Disclosure - Accumulated Other Comprehensive IncomeSheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureAccumulatedOtherComprehensiveIncomeAccumulated Other Comprehensive IncomeNotes17 falsefalseR18.htm995555 - Disclosure - Stock-Based CompensationSheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationStock-Based CompensationNotes18 falsefalseR19.htm995565 - Disclosure - WarrantsSheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureWarrantsWarrantsNotes19 falsefalseR20.htm995575 - Disclosure - Commitments and ContingenciesSheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureCommitmentsAndContingenciesCommitments and ContingenciesNotes20 falsefalseR21.htm995595 - Disclosure - Significant Accounting Policies (Policies)Sheethttp://nuvationbio.com/20240630/taxonomy/role/Role DisclosureSignificantAccountingPoliciesPoliciesSignificant Accounting Policies (Policies)PoliciesSheethttp://nuvationbio.com/20240630/taxonomy/role/Role DisclosureSignificantAccountingPolicies21 falsefalseR22.htm995605 - Disclosure - Significant Accounting Policies (Tables)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureSignificantAccountingPoliciesTablesSignificant Accounting Policies (Tables)TablesSheethttp://nuvationbio.com/20240630/taxonomy/role/Role DisclosureSignificantAccountingPolicies22 falsefalseR23.htm995615 - Disclosure - Acquisition (Tables)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureAcquisitionTablesAcquisition (Tables)TablesSheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureAcquisition23 falsefalseR24.htm995625 - Disclosure - Fair Value Measurements and Marketable Securities Available for Sale (Tables)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSaleTablesFair Value Measurements and Marketable Securities Available for Sale (Tables)TablesSheethttp://nuvationbio.com/20240630/taxonomy/role/Role DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSale24 falsefalseR25.htm995635 - Disclosure - Collaboration and License Agreements (Tables)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureCollaborationAndLicenseAgreementsTablesCollaboration and License Agreements (Tables)TablesSheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureCollaborationAndLicenseAgreements25 falsefalseR26.htm995645 - Disclosure - Leases (Tables)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureLeasesTablesLeases (Tables)TablesSheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureLeases26 falsefalseR27.htm995655 - Disclosure - Net Loss Per Share (Tables)Sheethttp://nuvationbio.com/20240630/taxonomy/role/Role DisclosureNetLossPerShareTablesNet Loss Per Share (Tables)TablesSheethttp://nuvationbio.com/20240630/taxonomy/role/Role DisclosureNetLossPerShare27 falsefalseR28.htm995665 - Disclosure - Accumulated Other Comprehensive Income (Tables)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureAccumulatedOtherComprehensiveIncomeTablesAccumulated Other Comprehensive Income (Tables)TablesSheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureAccumulatedOtherComprehensiveIncome28 falsefalseR29.htm995675 - Disclosure - Stock-Based Compensation (Tables)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationTablesStock-Based Compensation (Tables)TablesSheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensation29 falsefalseR30.htm995685 - Disclosure - Warrants (Tables)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureWarrantsTablesWarrants (Tables)TablesSheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureWarrants130 falsefalseR31.htm995705 - Disclosure - Significant Accounting Policies - Additional Information (Details)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureSignificantAccountingPoliciesAdditionalInformationDetailsSignificant Accounting Policies - Additional Information (Details)Details31 falsefalseR32.htm995715 - Disclosure - Significant Accounting Policies - Schedule of Concentration of Credit Risk Net Revenue (Details)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureSignificantAccountingPoliciesScheduleOfConcentrationOfCreditRiskNetRevenueDetailsSignificant Accounting Policies - Schedule of Concentration of Credit Risk Net Revenue (Details)Details32 falsefalseR33.htm995725 - Disclosure - Significant Accounting Policies - Schedule of Concentration of Customers and Suppliers Accounts Receivable (Details)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureSignificantAccountingPoliciesScheduleOfConcentrationOfCustomersAndSuppliersAccountsReceivable (Details)Details33 falsefalseR34.htm995745 - Disclosure - Acquisition - Additional Information (Details)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureAcquisitionAdditionalInformationDetailsAcquisition - Additional Information (Details)Details34 falsefalseR35.htm995755 - Disclosure - Acquisition - Summary of Final Allocable Purchase Price Consists (Details)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureAcquisitionSummaryOfFinalAllocablePurchasePriceConsistsDetailsAcquisition - Summary of Final Allocable Purchase Price Consists (Details)Details35 falsefalseR36.htm995765 - Disclosure - Acquisition - Schedule Of Final Allocable Purchase Price Consists (Parenthetical) (Details)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureAcquisitionScheduleOfFinalAllocablePurchasePriceConsistsParentheticalDetailsAcquisition - Schedule Of Final Allocable Purchase Price Consists (Parenthetical) (Details)Details36 falsefalseR37.htm995775 - Disclosure - Fair Value Measurements and Marketable Securities Available for Sale - Schedule of Marketable securities Measured at Fair Value on Recurring Basis (Details)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSaleScheduleOfMarketableSecuritiesMe Value Measurements and Marketable Securities Available for Sale - Schedule of Marketable securities Measured at Fair Value on Recurring Basis (Details)Details37 falsefalseR38.htm995785 - Disclosure - Fair Value Measurements and Marketable Securities Available for Sale - Schedule of Cost, Aggregate Fair Value and Unrealized Gains of Marketable Securities Available for sale (Details)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSaleScheduleOfCostAggregateFairValue Value Measurements and Marketable Securities Available for Sale - 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Disclosure - Collaboration and License Agreements - Schedule Of Contract Assets And Contract Liabilities (Details)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureCollaborationAndLicenseAgreementsScheduleOfContractAssetsAndContractLiabilitiesDetailsCollabora

[and License Agreements – Schedule Of Contract Assets And Contract Liabilities \(Details\)Details44 falsefalseR45.htm995865 – Disclosure – Borrowings – Additional Information](#)
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Payments (Details)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureLeasesScheduleOfAggregateLeasePaymentsDetailsLeases – Schedule Of Aggregate Lease
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(Details)Detailshttp://nuvationbio.com/20240630/taxonomy/role/Role-DisclosureNetLossPerShareSummaryOfSecuritiesOutstandingExcludedFromCalculationOfWeightedAverage
Loss Per Share – Summary of Securities Outstanding Excluded from Calculation of Weighted Average Shares Outstanding (Details)Details48 falsefalseR49.htm995905 –
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Option Activity under 2019 Plan and 2021 plan
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– Summary of Restricted Stock Unit Activity
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Values for Outstanding Warrants
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Hierarchy
(Details)Sheethttp://nuvationbio.com/20240630/taxonomy/role/DisclosureWarrantsSummaryOfChangesInLiabilitiesClassifiedInLevel3OfTheFairValueHierarchyDetailsWarrants
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Contingencies – Additional Information
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Additional Information (Details)Details58 falsefalseR59.htm996005 – Disclosure – Subsequent Event (Additional Information)
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Includes other kinds of accounts that have the general characteristics of demand deposits. Also includes short-term, highly liquid investments that are both readily convertible to known amounts of cash and so near their maturity that they present insignificant risk of changes in value because of changes in interest rates. 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Other information that may be disclosed includes (1) the nature of any restrictions on the entity's use of its cash and cash equivalents, (2) whether the entity's cash and cash equivalents are insured or expose the entity to credit risk, (3) the classification of any negative balance accounts (overdrafts), and (4) the carrying basis of cash equivalents (for example, at cost) and whether the carrying amount of cash equivalents approximates fair value." } } }, "auth ref": { "r30" } } }, "us-gaap_CashCashEquivalentsRestrictedCashAndRestrictedCashEquivalents": { "xbrltype": "monetaryItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "CashCashEquivalentsRestrictedCashAndRestrictedCashEquivalents", "crdr": "debit", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/StatementConsolidatedStatementsOfCashFlowsUnaudited": { "lang": { "en-us": { "role": { "periodStartLabel": "Cash and cash equivalents, beginning of the period", "periodEndLabel": "Cash and cash equivalents, end of the period", "label": "Cash, Cash Equivalents, Restricted Cash, and Restricted Cash Equivalents", "totalLabel": "Cash, Cash Equivalents, Restricted Cash and Restricted Cash Equivalents, Total", "documentation": "Amount of cash and cash equivalents, and cash and cash equivalents restricted to withdrawal or usage. Excludes amount for disposal group and discontinued operations. Cash includes, but is not limited to, currency on hand, demand deposits with banks or financial institutions, and other accounts with general characteristics of demand deposits. Cash equivalents include, but are not limited to, short-term, highly liquid investments that are both readily convertible to known amounts of cash and so near their maturity that they present insignificant risk of changes in value because of changes in interest rates." } } }, "auth ref": { "r29", "r89", "r201" } } }, "us-gaap_CashCashEquivalentsRestrictedCashAndRestrictedCashEquivalentsPeriodIncreaseDecreaseIncludingExchangeRateEffect": { "xbrltype": "monetaryItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "CashCashEquivalentsRestrictedCashAndRestrictedCashEquivalentsPeriodIncreaseDecreaseIncludingExchangeRateEffect", "crdr": "debit", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/StatementConsolidatedStatementsOfCashFlowsUnaudited": { "lang": { "en-us": { "role": { "totalLabel": "Net decrease in cash and cash equivalents", "label": "Cash, Cash Equivalents, Restricted Cash, and Restricted Cash Equivalents, Period Increase (Decrease), Including Exchange Rate Effect", "documentation": "Amount of increase (decrease) in cash, cash equivalents, and cash and cash equivalents restricted to withdrawal or usage, including effect from exchange rate change. Cash includes, but is not limited to, currency on hand, demand deposits with banks or financial institutions, and other accounts with general characteristics of demand deposits. 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Warrants and rights represent derivative securities that give the holder the right to purchase securities (usually equity) from the issuer at a specific price within a certain time frame. Warrants are often included in a new debt issue to entice investors by a higher return potential. The main difference between warrants and call options is that warrants are issued and guaranteed by the company, whereas options are exchange instruments and are not issued by the company. Also, the lifetime of a warrant is often measured in years, while the lifetime of a typical option is measured in months." } } }, "auth ref": "I1", "us-gaap ClassOfWarrantOrRightExercisePriceOfWarrantsOrRights": { "xbrltype": "perShareItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ClassOfWarrantOrRightExercisePriceOfWarrantsOrRights", "presentation": "http://nuvationbio.com/20240630/taxonomy/role/DisclosureSignificantAccountingPoliciesAdditionalInformationDetails", "http://nuvationbio.com/20240630/taxonomy/role/DisclosureSubsequentEventAdditionalInformationDetails", "lang": { "en-us": { "role": { "terseLabel": "Class of warrant or right, exercise price of warrants or rights", "label": "Class of Warrant or Right, Exercise Price of Warrants or Rights", "documentation": "Exercise price per share or per unit of warrants or rights outstanding." } } }, "auth ref": "r357", "us-gaap ClassOfWarrantOrRightLineItems": { "xbrltype": "stringItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ClassOfWarrantOrRightLineItems", "presentation": "http://nuvationbio.com/20240630/taxonomy/role/DisclosureWarrantsAdditionalInformationDetails", "http://nuvationbio.com/20240630/taxonomy/role/DisclosureWarrantsScheduleOffFairValuesForOutstandingWarrantsDetails", "http://nuvationbio.com/20240630/taxonomy/role/DisclosureWarrantsScheduleOffQuantitativeInformationRegardingLevel3FairValueMeasurementsDetails", "lang": { "en-us": { "role": { "terseLabel": "Class Of Warrant Or Right [Line Items]", "label": "Class of Warrant or Right [Line Items]", "documentation": "Line items represent financial concepts included in a table. 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The risk is the materially adverse effects of loss of a significant customer." } } }, "auth ref": { "r93", "r256" } } }, "nuvb DavidHanleyMember": { "xbrltype": "domainItemType", "nsuri": "http://nuvationbio.com/20240630", "localname": "DavidHanleyMember", "

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Includes beneficial interest in securitized financial asset." } }], "auth_ref": { "r126", "r306", "r765" } }, "us-gaap-DebtSecuritiesAvailableForSaleContinuousUnrealizedLossPositionLessThan12MonthsAccumulatedLoss": { "xbrltype": "monetaryItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "DebtSecuritiesAvailableForSaleContinuousUnrealizedLossPositionLessThan12MonthsAccumulatedLoss", "crdr": "debit", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSaleScheduleOfMarketableSecuritiesWithContinuous", "lang": { "en-us": { "role": { "label": "Debt Securities, Available for Sale, Continuous Unrealized Loss Position, Less than 12 Months, Accumulated Loss", "negatedLabel": "Less than 12 Months, Unrealized Losses", "documentation": "Amount of accumulated unrealized loss on investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available for sale), in continuous unrealized loss position for less than 12 months, without allowance for credit loss. Includes beneficial interest in securitized financial asset." } } }, "auth_ref": { "r126", "r306" } }, "us-gaap-DebtSecuritiesAvailableForSaleRealizedGain": { "xbrltype": "monetaryItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "DebtSecuritiesAvailableForSaleRealizedGain", "crdr": "credit", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSaleScheduleOfNetRealizedGainLossOnMarketableSe", "lang": { "en-us": { "role": { "label": "Debt Securities, Available for Sale, Realized Gain", "terseLabel": "Realized gains from sales of available-for-sale securities", "documentation": "Amount of realized gain on investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available for sale)." } } }, "auth_ref": { "r289" } }, "us-gaap-DebtSecuritiesAvailableForSaleRealizedLoss": { "xbrltype": "monetaryItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "DebtSecuritiesAvailableForSaleRealizedLoss", "crdr": "debit", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSaleScheduleOfNetRealizedGainLossOnMarketableSe", "lang": { "en-us": { "role": { "label": "Debt Securities, Available for Sale, Realized Loss", "negatedLabel": "Realized losses from sales of available-for-sale securities", "documentation": "Amount of realized loss on investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available for sale)." } } }, "auth_ref": { "r289" } }, "us-gaap-DebtSecuritiesAvailableForSaleUnrealizedGainLoss": { "xbrltype": "monetaryItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "DebtSecuritiesAvailableForSaleUnrealizedGainLoss", "crdr": "credit", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSaleScheduleOfNetRealizedGainLossOnMarketableSe", "lang": { "en-us": { "role": { "negatedLabel": "Amount reclassified for realized loss (gain) on marketable securities", "terseLabel": "Amount reclassified for realized loss (gain) on marketable securities", "totalLabel": "Debt Securities, Available for sale, Unrealized Gain (Loss), Total", "label": "Debt Securities, Available for Sale, Unrealized Gain (Loss)", "documentation": "Amount of unrealized gain (loss) on investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available for sale)." } } }, "auth_ref": { "r947", "r948" } }, "us-gaap-DebtSecuritiesAvailableForSaleUnrealizedLossPosition": { "xbrltype": "monetaryItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "DebtSecuritiesAvailableForSaleUnrealizedLossPosition", "crdr": "debit", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSaleScheduleOfMarketableSecuritiesWithContinuous", "lang": { "en-us": { "role": { "totalLabel": "Debt Securities, Available for sale, Unrealized Loss Position, Total", "label": "Debt Securities, Available for Sale, Unrealized Loss Position", "terseLabel": "Debt Securities, Available for sale, Unrealized Loss Position", "documentation": "Amount of investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available for sale), in unrealized loss position without allowance for credit loss." } } }, "auth_ref": { "r124", "r304", "r765" } }, "us-gaap-DebtSecuritiesAvailableForSaleUnrealizedLossPositionAccumulatedLoss": { "xbrltype": "monetaryItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "DebtSecuritiesAvailableForSaleUnrealizedLossPositionAccumulatedLoss", "crdr": "debit", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSaleScheduleOfMarketableSecuritiesWithContinuous", "lang": { "en-us": { "role": { "totalLabel": "Debt Securities, Available for sale, Unrealized Loss Position, Total", "label": "Debt Securities, Available for Sale, Unrealized Loss Position", "terseLabel": "Debt Securities, Available for sale, Unrealized Loss Position", "documentation": "Amount of investment in debt security measured at fair value with change in fair value recognized in other comprehensive income (available for sale), in unrealized loss position without allowance for credit loss. Includes beneficial interest in securitized financial asset." } } }, "auth_ref": { "r123", "r765", "r960" } }, "us-gaap-DebtSecuritiesAvailableforsaleAmortizedCostAbstract": { "xbrltype": "stringItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "DebtSecuritiesAvailableforsaleAmortizedCostAbstract", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSaleScheduleOfCostAggregateFairValueAndUnrealize", "lang": { "en-us": { "role": { "label": "Debt Securities, Available for Sale, Amortized Cost [Abstract]" } } }, "auth_ref": { " "us-gaap-DebtSecuritiesHeldtomaturityAllowanceForCreditLossLineItems": { "xbrltype": "stringItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "DebtSecuritiesHeldtomaturityAllowanceForCreditLossLineItems", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSaleScheduleOfMarketableSecuritiesWithContinuous", "lang": { "en-us": { "role": { "label": "Debt Securities, Held-to-Maturity, Allowance for Credit Loss [Line Items]", "documentation": "Line items represent financial concepts included in a table. 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Includes liabilities elected not to be offset. Excludes liabilities not subject to a master netting arrangement." } } }, "auth_ref": { "r18", "r76", "r107", "r108", "r185", "r754" } }, "us-gaap-DisaggregationOfRevenueTableTextBlock": { "xbrltype": "textBlockItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "DisaggregationOfRevenueTableTextBlock", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureCollaborationAndLicenseAgreementsTables", "lang": { "en-us": { "role": { "terseLabel": "Schedule of revenue recognized", "label": "Disaggregation of Revenue [Table Text Block]", "documentation": "Tabular disclosure of disaggregation of revenue into categories depicting how nature, amount, timing, and uncertainty of revenue and cash flows are affected by economic factor." } } }, "auth_ref": { "r975" } }, "us-gaap-DisclosureOfCompensationRelatedCostsShareBasedPaymentsTextBlock": { "xbrltype": "textBlockItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "DisclosureOfCompensationRelatedCostsShareBasedPaymentsTextBlock", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensation", "lang": { "en-us": { "role": { "label": "Share-Based Payment Arrangement [Text Block]", "terseLabel": "Stock-Based Compensation", "documentation": "The entire disclosure for share-based payment arrangement." } } }, "auth_ref": { "r381", "r385", "r416", "r417", "r418", "r778" } }, "us-gaap-DisclosureOfCompensationRelatedCostsShareBasedPaymentsAbstract": { "xbrltype": "stringItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "DisclosureOfCompensationRelatedCostsShareBasedPaymentsAbstract", "lang": { "en-us": { "role": { "label": "Share-Based Payment Arrangement [Abstract]" } } }, "auth_ref": { " "dei DocumentFiscalPeriodFocus": { "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DocumentDocumentAndEntityInformation", "lang": { "en-us": { "role": { "terseLabel": "Document Fiscal Period Focus", "label": "Document Fiscal Period Focus", "documentation": "Fiscal period values are FY, Q1, Q2, and Q3. 1st, 2nd and 3rd quarter 10-Q or 10-QT statements have value Q1, Q2, and Q3 respectively, with 10-K, 10-KT or other fiscal year statements having FY." } } }, "auth_ref": { " "dei DocumentFiscalYearFocus": { "xbrltype": "yearItemType", "nsuri": "http://xbrl.sec.gov/dei/2024", "localname": "DocumentFiscalYearFocus", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DocumentDocumentAndEntityInformation", "lang": { "en-us": { "role": { "terseLabel": "Document Fiscal Year Focus", "label": "Document Fiscal Year Focus", "documentation": "This is focus fiscal year of the document report in YYYY format. 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Excludes amounts for disposal group and discontinued operations. Cash includes, but is not limited to, currency on hand, demand deposits with banks or financial institutions, and other accounts with general characteristics of demand deposits. Cash equivalents include, but are not limited to, short-term, highly liquid investments that are both readily convertible to known amounts of cash and so near their maturity that they present insignificant risk of changes in value because of changes in interest rates." } } }, "auth ref": { "r488": { "us-gaap EmployeeServiceShareBasedCompensationAllocationOfRecognizedPeriodCostsLineItems": { "xbrltype": "stringItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "EmployeeServiceShareBasedCompensationAllocationOfRecognizedPeriodCostsLineItems", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationSummaryOfRestrictedStockUnitActivityDetails": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationSummaryOfStockOptionActivityUnder2019PlanAnd2021PlanDetails": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationSummaryOfStockbasedCompensationExpenseDetails": { "lang": { "en-us": { "role": { "terseLabel": "Employee Service Share Based Compensation Allocation Of Recognized Period Costs [Line Items]", "label": "Share-Based Payment Arrangement, Expensed and Capitalized, Amount [Line Items]", "documentation": "Line items represent financial concepts included in a table. 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Where multiple classes or units exist define each class/interest by adding class of stock items such as Common Class A [Member], Common Class B [Member] or Partnership Interest [Member] onto the Instrument [Domain] of the Entity Listings, Instrument." } } }, "auth ref": { "r219": { "dei EntityCurrentReportingStatus": { "xbrltype": "yesNoItemType", "nsuri": "http://xbrl.sec.gov/dei/2024", "localname": "EntityCurrentReportingStatus", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DocumentDocumentAndEntityInformation": { "lang": { "en-us": { "role": { "terseLabel": "Entity Current Reporting Status", "label": "Entity Current Reporting Status", "documentation": "Indicate 'Yes' or 'No' whether registrants (1) have filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months for for such shorter period that registrants were required to file such reports; and (2) have been subject to such filing requirements for the past 90 days. 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The disclosures contemplated herein include the fair value measurements at the reporting date by the level within the fair value hierarchy in which the fair value measurements in their entirety fall, segregating fair value measurements using quoted prices in active markets for identical assets (Level 1), significant other observable inputs (Level 2), and significant unobservable inputs (Level 3). 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Examples of items that might be included in the application of this element may consist of letters of credit, standby letters of credit, and revolving credit arrangements, under which borrowings can be made up to a maximum amount as of any point in time conditional on satisfaction of specified terms before, as of and after the date of drawdowns on the line. Includes short-term obligations that would normally be classified as current liabilities but for which (a) postbalance sheet date issuance of a long-term obligation to refinance the short-term obligation on a long-term basis, or (b) the enterprise has entered into a financing agreement that clearly permits the enterprise to refinance the short-term obligation on a long-term basis and the following conditions are met (1) the agreement does not expire within 1 year and is not cancelable by the lender except for violation of an objectively determinable provision, (2) no violation exists at the BS date, and (3) the lender has entered into the financing agreement is expected to be financially capable of honoring the agreement.", "auth ref": "r20", "r112", "r1032", "us-gaap LineOfCreditFacilityFrequencyOfCommitmentFeePayment", "xbrltype": "stringItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "LineOfCreditFacilityFrequencyOfCommitmentFeePayment", "presentation": 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[illegible]

"Nonrecurring Adjustment [Axis]", "documentation": "Information by nonrecurring adjustment directly attributable to business combination included in pro forma revenue and earnings." } } }, "auth ref": { "r16" } }, "us-gaap-NonrecurringAdjustmentDomain": { "xbrltype": "domainItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "NonrecurringAdjustmentDomain", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureAcquisitionSummaryOfFinalAllocablePurchasePriceConsistsDetails": { "lang": { "en-us": { "role": { "label": "Nonrecurring Adjustment [Domain]", "documentation": "Material, nonrecurring adjustment(s) allocated (included) to (in) reported pro forma revenue and earnings (supplemental pro forma information)." } } }, "auth ref": { "r16" } }, "nuvb-NumberOfEquitySecuritiesSold": { "xbrltype": "integerItemType", "nsuri": "http://nuvationbio.com/20240630", "localname": "NumberOfEquitySecuritiesSold", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSaleAdditionalInformationDetails": { "lang": { "en-us": { "role": { "documentation": "Number of Equity Securities Sold.", "label": "Number of Equity Securities Sold", "terseLabel": "Number of equity securities sold" } } }, "auth ref": { "nuvb-NumberOfSharesIssuableUnderShareBasedPaymentArrangement": { "xbrltype": "sharesItemType", "nsuri": "http://nuvationbio.com/20240630", "localname": "NumberOfSharesIssuableUnderShareBasedPaymentArrangement", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationAdditionalInformationDetails": { "lang": { "en-us": { "role": { "terseLabel": "Number of common shares issuable under the plan (in shares)", "label": "Number Of Shares Issuable Under Share Based Payment Arrangement", "documentation": "Number of shares issuable under share based payment arrangement." } } }, "auth ref": { "us-gaap-OperatingExpenses": { "xbrltype": "monetaryItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "OperatingExpenses", "crdr": "debit", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/StatementConsolidatedStatementsOfOperationsAndComprehensiveLossUnaudited": { "lang": { "en-us": { "role": { "totalLabel": "Total operating expenses", "label": "Operating Expenses", "documentation": "Generally recurring costs associated with normal operations except for the portion of these expenses which can be clearly related to production and included in cost of sales or services. Includes selling, general and administrative expense." } } }, "auth ref": { "us-gaap-OperatingExpensesAbstract": { "xbrltype": "stringItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "OperatingExpensesAbstract", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/StatementConsolidatedStatementsOfOperationsAndComprehensiveLossUnaudited": { "lang": { "en-us": { "role": { "terseLabel": "Operating expenses", "label": "Operating Expenses [Abstract]" } } }, "auth ref": { "us-gaap-OperatingIncomeLoss": { "xbrltype": "monetaryItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "OperatingIncomeLoss", "crdr": "credit", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/StatementConsolidatedStatementsOfOperationsAndComprehensiveLossUnaudited": { "lang": { "en-us": { "role": { "terseLabel": "Loss from operations", "label": "Operating Income (Loss)", "totalLabel": "Loss from operations", "documentation": "The net result for the period of deducting operating expenses from operating revenues." } } }, "auth ref": { "r121", "r758", "r940", "r941", "r942", "r943", "r944" } }, "us-gaap-OperatingLeaseCost": { "xbrltype": "monetaryItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "OperatingLeaseCost", "crdr": "debit", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureLeasesAdditionalInformationDetails": { "lang": { "en-us": { "role": { "label": "Operating Lease, Cost", "terseLabel": "Operating Leases", "documentation": "Amount of single lease cost, calculated by allocation of remaining cost of lease over remaining lease term. 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Note: elements for number of nonredeemable preferred shares, par value and other disclosure concepts are in another section within stockholders' equity." } } }, "auth ref": { "r70", "r595", "r793" } }, "us-gaap PrepaidExpenseAndOtherAssetsCurrent": { "xbrltype": "monetaryItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "PrepaidExpenseAndOtherAssetsCurrent", "crdr": "debit", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/StatementConsolidatedBalanceSheetsUnaudited": { "lang": { "en-us": { "role": { "terseLabel": "Prepaid expenses and other current assets", "label": "Prepaid Expense and Other Assets, Current", "documentation": "Amount of asset related to consideration paid in advance for costs that provide economic benefits in future periods, and amount of other assets that are expected to be realized or consumed within one year or the normal operating cycle, if longer." } } }, "auth ref": { "r925" } }, "nuvb PrivatePlacementWarrantsMember": { 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"http://nuvationbio.com/20240630/taxonomy/role/Role_DisclosureCollaborationAndLicenseAgreementsScheduleOfRevenueRecognizedDetails",
"http://nuvationbio.com/20240630/taxonomy/role/StatementConsolidatedStatementsOfOperationsAndComprehensiveIncomeLossUnaudited"} l": "lang", { "en-us": { "role": {
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authority that is both imposed on and concurrent with specific revenue-producing transaction, including, but not limited to, sales, use, value added and excise." } },
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Stock, Description of Transaction", "label": "Sale of Stock, Description of Transaction", "documentation": "Description of stock transaction which may include details of the
offering (IPO, private placement), a description of the stock sold, percentage of subsidiary's or equity investee's stock sold, a description of the investors and whether the
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Includes, but is not limited to, recognized asset and liability." } } }, "auth ref": { "r51", "r53", "r443" } } }, "us-gaap:ScheduleOfBusinessAcquisitionsByAcquisitionTextBlock": { "xbrltype": "textBlockItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ScheduleOfBusinessAcquisitionsByAcquisitionTextBlock", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureAcquisitionTables", "lang": { "en-us": { "role": { "terseLabel": "Schedule of Final Allocable Purchase Price Consists", "label": "Schedule of Business Acquisitions, by Acquisition [Table Text Block]", "documentation": "Tabular disclosure of a material business combination completed during the period, including background, timing, and recognized assets and liabilities. This table does not include leveraged buyouts." } } }, "auth ref": { "r51", "r53" } } }, "us-gaap:ScheduleOfDefinedBenefitPlansDisclosuresTable": { "xbrltype": "stringItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ScheduleOfDefinedBenefitPlansDisclosuresTable", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableForSaleScheduleOfMarketableSecuritiesMeasuredAtFair", "lang": { "en-us": { "role": { "terseLabel": "Schedule Of Defined Benefit Plans Disclosures [Table]", "label": "Defined Benefit Plan [Table]", "documentation": "Disclosure of information about individual defined benefit pension plan or other postretirement defined benefit plan. It may be appropriate to group certain similar plans. Also includes schedule for fair value of plan assets by major categories of plan assets by the level within the fair value hierarchy in which the fair value measurements in their entirety fall, segregating fair value measurements using quoted prices in active markets for identical assets or liabilities (Level 1), Significant other observable inputs (Level 2), and significant unobservable inputs (Level 3)." } } }, "auth ref": { "r10", "r42", "r43", "r44", "r45" } } }, "us-gaap:ScheduleOfEmployeeServiceShareBasedCompensationAllocationOfRecognizedPeriodCostsTable": { "xbrltype": "stringItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ScheduleOfEmployeeServiceShareBasedCompensationAllocationOfRecognizedPeriodCostsTable", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationSummaryOfRestrictedStockUnitActivityDetails", "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationSummaryOfStockOptionActivityUnder2019PlanAnd2021PlanDetails", "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationSummaryOfStockbasedCompensationExpenseDetails", "lang": { "en-us": { "role": { "terseLabel": "Schedule Of Employee Service Share Based Compensation Allocation Of Recognized Period Costs [Table]", "label": "Share-Based Payment Arrangement, Expensed and Capitalized, Amount [Table]", "documentation": "Disclosure of information about amount recognized for award under share-based payment arrangement. Includes, but is not limited to, amount expensed in statement of income or comprehensive income, amount capitalized in statement of financial position, and corresponding reporting line item in financial statements." } } }, "auth ref": { "r48" } } }, "us-gaap:ScheduleOfEmployeeServiceShareBasedCompensationAllocationOfRecognizedPeriodCostsTextBlock": { "xbrltype": "textBlockItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ScheduleOfEmployeeServiceShareBasedCompensationAllocationOfRecognizedPeriodCostsTextBlock", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationTables", "lang": { "en-us": { "role": { "label": "Share-Based Payment Arrangement, Expensed and Capitalized, Amount [Table Text Block]", "terseLabel": "Summary of Stock-Based Compensation Expense", "documentation": "Tabular disclosure of allocation of amount expensed and capitalized for award under share-based payment arrangement to statement of income or comprehensive income and statement of financial position. Includes, but is not limited to, corresponding line item in financial statement." } } }, "auth ref": { "r48" } } }, "us-gaap:ScheduleOfRealizedGainLossTableTextBlock": { "xbrltype": "textBlockItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ScheduleOfRealizedGainLossTableTextBlock", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureFairValueMeasurementsAndMarketableSecuritiesAvailableforsaleTables", "lang": { "en-us": { "role": { "label": "Schedule of Realized Gain (Loss) [Table Text Block]", "terseLabel": "Schedule of Net Realized Gain (Loss) on Marketable Securities", "documentation": "Tabular disclosure of the proceeds from sales of available-for-sale securities and the gross realized gains and gross realized losses that have been included in earnings as a result of those sales." } } }, "auth ref": { "r953" } } }, "us-gaap:ScheduleOfShareBasedCompensationArrangementsByShareBasedPaymentAwardTable": { "xbrltype": "stringItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ScheduleOfShareBasedCompensationArrangementsByShareBasedPaymentAwardTable", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationAdditionalInformationDetails", "lang": { "en-us": { "role": { "terseLabel": "Schedule Of Share Based Compensation Arrangements By Share Based Payment Award [Table]", "label": "Schedule of Share-Based Compensation Arrangements by Share-Based Payment Award [Table]", "documentation": "Disclosure of information about share-based payment arrangement." } } }, "auth ref": { "r382", "r384", "r386", "r387", "r388", "r390", "r391", "r392", "r393", "r394", "r395", "r396", "r397", "r398", "r399", "r400", "r401", "r402", "r403", "r404", "r405", "r406", "r407", "r410", "r411", "r412", "r413", "r414" } } }, "us-gaap:ScheduleOfShareBasedCompensationRestrictedStockUnitsAwardActivityTableTextBlock": { "xbrltype": "textBlockItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ScheduleOfShareBasedCompensationRestrictedStockUnitsAwardActivityTableTextBlock", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationTables", "lang": { "en-us": { "role": { "terseLabel": "Summary of Restricted Stock Units Activity", "label": "Share-Based Payment Arrangement, Restricted Stock Unit, Activity [Table Text Block]", "documentation": "Tabular disclosure of the number and weighted-average grant date fair value for restricted stock units that were outstanding at the beginning and end of the year, and the number of restricted stock units that were granted, vested, or forfeited during the year." } } }, "auth ref": { "r102" } } }, "us-gaap:ScheduleOfShareBasedCompensationStockOptionsActivityTableTextBlock": { "xbrltype": "textBlockItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ScheduleOfShareBasedCompensationStockOptionsActivityTableTextBlock", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationTables", "lang": { "en-us": { "role": { "label": "Share-Based Payment Arrangement, Option, Activity [Table Text Block]", "terseLabel": "Summary of Stock Option Activity under 2019 Plan and 2021 Plan", "documentation": "Tabular disclosure for stock option plans. Includes, but is not limited to, outstanding awards at beginning and end of year, grants, exercises, forfeitures, and weighted-average grant date fair value." } } }, "auth ref": { "r14", "r15", "r102" } } }, "us-gaap:ScheduleOfStockholdersEquityNoteWarrantsOrRightsTextBlock": { "xbrltype": "textBlockItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ScheduleOfStockholdersEquityNoteWarrantsOrRightsTextBlock", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureWarrantsTables", "lang": { "en-us": { "role": { "label": "Schedule of Stockholders' Equity Note, Warrants or Rights [Table Text Block]", "terseLabel": "Schedule of Fair Values for Outstanding Warrants", "documentation": "Tabular disclosure of warrants or rights issued. Warrants and rights outstanding are derivative securities that give the holder the right to purchase securities (usually equity) from the issuer at a specific price within a certain time frame. Warrants are often included in a new debt issue to entice investors by a higher return potential. The main difference between warrants and call options is that warrants are issued and guaranteed by the company, whereas options are exchange instruments and are not issued by the company. Also, the lifetime of a warrant is often measured in years, while the lifetime of a typical option is measured in months. Disclose the title of issue of securities called for by warrants and rights outstanding, the aggregate amount of securities called for by warrants and rights outstanding, the date from which the warrants or rights are exercisable, and the price at which the warrant or right is exercisable." } } }, "auth ref": { "r46" } } }, "us-gaap:SchedulesOfConcentrationOfRiskByRiskFactorTextBlock": { "xbrltype": "textBlockItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "SchedulesOfConcentrationOfRiskByRiskFactorTextBlock", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureSignificantAccountingPoliciesTables", "lang": { "en-us": { "role": { "terseLabel": "Schedule of Concentration of Risk Concentration of customers", "label": "Schedules of Concentration of Risk, by Risk Factor [Table Text Block]", "documentation": "Tabular disclosure of the nature of a concentration, a benchmark to which it is compared, and the percentage that the risk is to the benchmark." } } }, "auth ref": { "r32", "r33", "r34", "r35", "r57", "r109" } } }, "dei:Security12bTitle": { "xbrltype": "securityTitleItemType", "nsuri": "http://xbrl.sec.gov/dei/2024", "localname": "Security12bTitle", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DocumentDocumentAndEntityInformation", "lang": { "en-us": { "role": { "terseLabel": "Title of 12(b) Security", "label": "Title of 12(b) Security", "documentation": "Title of a 12(b) registered security." } } }, "auth ref": { "r821" } } }, "dei:SecurityExchangeName": { "xbrltype": "securityExchangeName", "nsuri": "http://xbrl.sec.gov/dei/2024", "localname": "SecurityExchangeName", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DocumentDocumentAndEntityInformation", "lang": { "en-us": { "role": { "terseLabel": "Security Exchange Name", "label": "Security Exchange Name", "documentation": "Name of the Exchange on which a security is registered." } } }, "auth ref": { "r823" } } }, "srt:SegmentGeographicalDomain": { "xbrltype": "domainItemType", "nsuri": "http://fasb.org/srt/2024", "localname": "SegmentGeographicalDomain", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureLeasesAdditionalInformationDetails", "http://nuvationbio.com/20240630/taxonomy/role/DisclosureSignificantAccountingPoliciesAdditionalInformationDetails", "lang": { "en-us": { "role": { "label": "Geographical Domain", "documentation": "Geographical area." } } }, "auth ref": { "r144", "r146", "r149", "r151", "r158", "r159", "r160", "r161", "r162", "r163", "r164", "r165", "r166", "r167", "r254", "r255", "r554", "r555", "r556", "r557", "r558", "r559", "r560", "r561", "r562", "r563", "r564", "r565", "r566", "r567", "r568", "r569", "r570", "r571", "r572", "r573", "r574", "r575", "r576", "r577", "r578", "r579", "r580", "r581", "r582", "r583", "r658", "r659", "r660", "r668", "r678", "r722", "r725", "r727", "r734", "r738", "r739", "r740", "r741", "r742", "r743", "r744", "r745", "r746", "r752", "r771", "r795", "r796", "r797", "r798", "r799", "r800", "r801", "r802", "r806", "r812", "r975", "r1036", "r1038", "r1039", "r1041", "r1042", "r1043", "r1044", "r1045", "r1046", "r1047", "r1048", "r1049", "r1050", "r1051", "r1052", "r1053", "r1054", "r1055", "r1056", "r1057", "r1058", "r1059", "r1060", "r1061", "r1062", "r1063", "r1065", "r1066" } } }, "us-gaap:SegmentReportingPolicyPolicyTextBlock": { "xbrltype": "textBlockItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "SegmentReportingPolicyPolicyTextBlock", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/Role-DisclosureSignificantAccountingPoliciesPolicies", "lang": { "en-us": { "role": { "terseLabel": "Segment Information", "label": "Segment Reporting, Policy [Policy Text Block]", "documentation": "Disclosure of accounting policy for segment reporting." } } }, "auth ref": { "r242", "r243", "r244", "r245", "r246", "r247", "r248", "r252", "r254", "r259", "r760", "r763" } } }, "us-gaap:SellingGeneralAndAdministrativeExpense": { "xbrltype": "monetaryItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "SellingGeneralAndAdministrativeExpense", "crdr": "debit", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/StatementConsolidatedStatementsOfOperationsAndComprehensiveLossUnaudited", "lang": { "en-us": { "role": { "terseLabel": "Selling, general and administrative", "label": "Selling, General and Administrative Expense", "totalLabel": "Selling, General and Administrative Expense, Total", "documentation": "The aggregate total costs related to selling a firm's product and services, as well as all other general and administrative expenses. Direct selling expenses (for example, credit, warranty, and advertising) are expenses that can be directly linked to the sale of specific products. Indirect selling expenses are expenses that cannot be directly linked to the sale of specific products, for example telephone expenses, Internet, and postal charges. General and administrative expenses include salaries of non-sales personnel, rent, utilities, communication, etc." } } }, "auth ref": { "r84" } } }, "us-gaap:SellingGeneralAndAdministrativeExpensesMember": { "xbrltype": "domainItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "SellingGeneralAndAdministrativeExpensesMember", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationSummaryOfStockbasedCompensationExpenseDetails", "lang": { "en-us": { "role": { "label": "Selling, General and Administrative Expenses [Member]", "terseLabel": "General and Administrative [Member]", "documentation": "Primary financial statement caption encompassing selling, general and administrative expense." } } }, "auth ref": { "r1" } } }, "us-gaap:SeriesAPreferredStockMember": { "xbrltype": "domainItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "SeriesAPreferredStockMember", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureAcquisitionAdditionalInformationDetails", "lang": { "en-us": { "role": { "terseLabel": "Series A Preferred Stock", "label": "Series A Preferred Stock [Member]", "documentation": "Series A preferred stock." } } }, "auth ref": { "r923", "r924", "r973" } } }, "us-gaap:ShareBasedCompensation": { "xbrltype": "monetaryItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ShareBasedCompensation", "crdr": "debit", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/StatementConsolidatedStatementsOfCashFlowsUnaudited", "lang": { "en-us": { "role": { "terseLabel": "Stock-based compensation", "label": "Share-Based Payment Arrangement, Noncash Expense", "totalLabel": "Share-based Payment Arrangement, Noncash Expense, Total", "documentation": "Amount of noncash expense for share-based payment arrangement." } } }, "auth ref": { "r77" } } }, "us-gaap:ShareBasedCompensationArrangementByShareBasedPaymentAwardDescription": { "xbrltype": "stringItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ShareBasedCompensationArrangementByShareBasedPaymentAwardDescription", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationAdditionalInformationDetails", "lang": { "en-us": { "role": { "terseLabel": "Share-based compensation arrangement by share-based payment award, description", "label": "Share-Based Compensation Arrangement by Share-Based Payment Award, Description", "documentation": "Description of terms of share-based payment arrangement. Includes, but is not limited to, type of award or grantee and reason for issuance." } } }, "auth ref": { "r46", "r47" } } }, "us-gaap:ShareBasedCompensationArrangementByShareBasedPaymentAwardFairValueAssumptionsExpectedDividendRate": { "xbrltype": "percentItemType", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ShareBasedCompensationArrangementByShareBasedPaymentAwardFairValueAssumptionsExpectedDividendRate", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureWarrantsScheduleOfQuantitativeInformationRegardingLevel3FairValueMeasurementsDetails", "lang": { "en-us":

{ "role": { "label": "Share-based Compensation Arrangement by Share-based Payment Award, Fair Value Assumptions, Expected Dividend Rate", "terseLabel": "Dividend", "documentation": "The estimated dividend rate (a percentage of the share price) to be paid (expected dividends) to holders of the underlying shares over the option's term." } }, { "auth_ref": { "r412" } }, { "us-gaap_ShareBasedCompensationArrangementByShareBasedPaymentAwardFairValueAssumptionsExpectedVolatilityRate": { "xbrltype": "percentItem", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ShareBasedCompensationArrangementByShareBasedPaymentAwardFairValueAssumptionsExpectedVolatilityRate", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureWarrantsScheduleOfQuantitativeInformationRegardingLevel3FairValueMeasurementsDetails" } }, { "lang": { "en-us": { "role": { "label": "Share-based Compensation Arrangement by Share-based Payment Award, Fair Value Assumptions, Expected Volatility Rate", "terseLabel": "Expected volatility", "documentation": "The estimated measure of the percentage by which a share price is expected to fluctuate during a period. Volatility also may be defined as a probability-weighted measure of the dispersion of returns about the mean. The volatility of a share price is the standard deviation of the continuously compounded rates of return on the share over a specified period. That is the same as the standard deviation of the differences in the natural logarithms of the stock prices plus dividends, if any, over the period." } } }, { "auth_ref": { "r411" } }, { "us-gaap_ShareBasedCompensationArrangementByShareBasedPaymentAwardFairValueAssumptionsRiskFreeInterestRate": { "xbrltype": "percentItem", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ShareBasedCompensationArrangementByShareBasedPaymentAwardFairValueAssumptionsRiskFreeInterestRate", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureWarrantsScheduleOfQuantitativeInformationRegardingLevel3FairValueMeasurementsDetails" } }, { "lang": { "en-us": { "role": { "label": "Share-based Compensation Arrangement by Share-based Payment Award, Fair Value Assumptions, Risk Free Interest Rate", "terseLabel": "Risk-free interest rate", "documentation": "The risk-free interest rate assumption that is used in valuing an option on its own shares." } } }, { "auth_ref": { "r413" } }, { "us-gaap_ShareBasedCompensationArrangementByShareBasedPaymentAwardLineItems": { "xbrltype": "stringItem", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ShareBasedCompensationArrangementByShareBasedPaymentAwardLineItems", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationAdditionalInformationDetails" } }, { "lang": { "en-us": { "role": { "label": "Share-Based Compensation Arrangement by Share-Based Payment Award [Line Items]", "label": "Share-Based Compensation Arrangement by Share-Based Payment Award [Line Items]", "documentation": "Line items represent financial concepts included in a table. These concepts are used to disclose reportable information associated with domain members defined in one or many axes to the table." } } }, { "auth_ref": { "r382", "r384", "r386", "r387", "r388", "r390", "r391", "r392", "r393", "r394", "r395", "r396", "r397", "r398", "r399", "r400", "r401", "r402", "r403", "r404", "r405", "r406", "r407", "r410", "r411", "r412", "r413", "r414" } }, { "us-gaap_ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsExercisableNumber": { "xbrltype": "sharesItem", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsExercisableNumber", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationSummaryOfStockOptionActivityUnder2019PlanAnd2021PlanDetails" } }, { "lang": { "en-us": { "role": { "terseLabel": "Exercisable at June 30, 2024", "label": "Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Exercisable, Number", "documentation": "The number of shares into which fully or partially vested stock options outstanding as of the balance sheet date can be currently converted under the option plan." } } }, { "auth_ref": { "r392" } }, { "us-gaap_ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsForfeituresInPeriod": { "xbrltype": "sharesItem", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsForfeituresInPeriod", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationSummaryOfStockOptionActivityUnder2019PlanAnd2021PlanDetails" } }, { "lang": { "en-us": { "role": { "label": "Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Forfeitures in Period", "terseLabel": "Number of share option, Forfeited", "negatedLabel": "Forfeited", "documentation": "The number of shares under options that were cancelled during the reporting period as a result of occurrence of a terminating event specified in contractual agreements pertaining to the stock option plan." } } }, { "auth_ref": { "r396" } }, { "us-gaap_ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsGrantsInPeriodGross": { "xbrltype": "sharesItem", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsGrantsInPeriodGross", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationSummaryOfRestrictedStockUnitActivityDetails" } }, { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationSummaryOfStockOptionActivityUnder2019PlanAnd2021PlanDetails" } }, { "lang": { "en-us": { "role": { "label": "Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Grants in Period, Gross", "verboseLabel": "Granted", "documentation": "Gross number of share options (or share units) granted during the period." } } }, { "auth_ref": { "r394" } }, { "us-gaap_ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsOutstandingNumber": { "xbrltype": "sharesItem", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsOutstandingNumber", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationSummaryOfStockOptionActivityUnder2019PlanAnd2021PlanDetails" } }, { "lang": { "en-us": { "role": { "periodEndLabel": "Outstanding at June 30, 2024", "label": "Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Outstanding, Number", "periodStartLabel": "Outstanding at December 31, 2023", "documentation": "Number of options outstanding, including both vested and non-vested options." } } }, { "auth_ref": { "r390", "r391" } }, { "us-gaap_ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsOutstandingPeriodIncreaseDecreaseWeightedAverageExercisePrice": { "xbrltype": "perShareItem", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ShareBasedCompensationArrangementByShareBasedPaymentAwardOptionsOutstandingPeriodIncreaseDecreaseWeightedAverageExercisePrice", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureStockbasedCompensationAdditionalInformationDetails" } }, { "lang": { "en-us": { "role": { "label": "Share-Based Compensation Arrangement by Share-Based Payment Award, Options, Outstanding, Period Increase (Decrease), Weighted Average Exercise Price", "terseLabel": "Weighted average exercise price of outstanding options", "documentation": "Change in the weighted average exercise price of options outstanding." } } }, { "auth_ref": { "r381", "r382", "r383", "r384", "r385", "r386", "r387", "r388", "r389", "r390", "r391", "r392", "r393", "r394", "r395", "r396", "r397", "r398", "r399", "r400", "r401", "r402", "r403", "r404", "r405", "r406", "r407", "r410", "r411", "r412", "r413", "r414" } }, { "us-gaap_ShareBasedCompensationOptionAndIncentivePlansPolicy": { "xbrltype": "textBlockItem", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "ShareBasedCompensationOptionAndIncentivePlansPolicy", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/Role-DisclosureSignificantAccountingPoliciesPolicies" } }, { "lang": { "en-us": { "role": { "terseLabel": "Stock-based Compensation", "label": "Share-Based Payment Arrangement [Policy Text Block]", "documentation": "Disclosure of accounting policy for award under share-based payment arrangement. Includes, but is not limited to, methodology and assumption used in measuring cost." } } }, { "auth_ref": { "r381", "r389", "r408", "r409", "r410", "r411", "r414", "r419", "r420", "r421", "r422" } }, { "us-gaap_SharePrice": { "xbrltype": "perShareItem", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "SharePrice", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/DisclosureWarrantsScheduleOfQuantitativeInformationRegardingLevel3FairValueMeasurementsDetails" } }, { "lang": { "en-us": { "role": { "label": "Share Price", "documentation": "Price of a single share of a number of saleable stocks of a company." } } }, { "auth_ref": { "r410" } }, { "us-gaap_SharebasedCompensationArrangementBySharebasedPaymentAwardFairValueAssumptionsExpectedTerm": { "xbrltype": "durationItem", "nsuri": "http://fasb.org/us-gaap/2024", "localname": "SharebasedCompensationArrangementBySharebasedPaymentAwardFairValueAssumptionsExpectedTerm", "presentation": { 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Such securities may include treasury bills (short-term maturities - one year or less), treasury notes (intermediate term maturities - two to ten years), and treasury bonds (long-term maturities - ten to thirty years), debt securities issued by the Government National Mortgage Association (Ginnie Mae) and debt securities issued by the Federal National Mortgage Association (Fannie Mae) or the Federal Home Loan Mortgage Corporation (Freddie Mac)." } } } , "auth ref": { "r590", "r772", "r794", "r808", "r1064" } } , "nuvb UnauditedInterimFinancialInformationPolicyTextBlock": { "xbrltype": "textBlockItemType", "nsuri": "http://nuvationbio.com/20240630", "localname": "UnauditedInterimFinancialInformationPolicyTextBlock", "presentation": { "http://nuvationbio.com/20240630/taxonomy/role/Role-DisclosureSignificantAccountingPoliciesPolicies" } , "lang": { "en-us": { "role": { "terseLabel": "Unaudited Interim Financial Information", "label": "Unaudited Interim Financial Information Policy [Text 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M99@<16P7J.YM.6PCL=885D)BNJ2D-QFG\$E+H.LJ3U7628HQH!H!<1+9?7.MD3.N6G!A.LX+!H&HMP9J2IYW\$!<JCYAKV!<0&SK272.0W!8FUM&6P.M.65RV?>
O&L!<=<KMMU7IWSA~!541KY4BVA!<H&RPN>1<=<U&A5J5V.M?<F!<ONZ+<R2QD?T4E5M!G2H\$F5.S.2N\$8EB8!0VYI(L!<90%~.MEL#8672E?2JQ4TOH5FFD.A8~6WY.T3+!<!
100L.P&VY!<F.7+!A.ME!JIM&5<6%L.RW=<=<KSI(D.MY~2~!K.L3.2DU@8KJ!>37<~M.S.POX.MPY.+!XJ2P.DH4Y59=F&D8HJQD!JFFH&RQD!GOD(NB!+I9X1!<H&@5
M.%!<=<8F<=<1\$<G~NM.L.9.74!<T!ED9X8?>!JW7260E!D.YV!+<~W03@%<MXP!H!BK!WIG62~\$<D+OQ>IV26LIG0EQZB&Q@<~!V:0MF=SQ=<AUN8?7!<=<
M(50BLG<9P0B9=H!+!WGTV7!W!P&FAYQ75SWW?N.DY.319~NBR+M@T%<M.UX-Y2ZL&F&W4P56D=XJ.C.7ABX~!D.T.9.RT>=<Q.7&L@!<X&E6B2DGSUW.MB!<4*=<
Y2ZRP!<L05J&GR31>H!<T&SOQ-YJ2Q7ZBWJ.YI.G2+!<SBL\$U8!D!M015X&M7B18U!<IOM<K."SAI~0*0*SYK~<K&B54\$B!5!<X\$=<N2W\$5~.M63NWB80?F4I!C3R.7>~!HET
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M@<@<4!<X>=>~PB927W6I7T074.62>~<V2.R@K!P!HXW8&.<C!<OH\$<BW~M8!Y.LD2W1G!<0D?>HS(YGCS(Q!<L.M.GODDKWZ!<=<O&H=53Z%YR!N!<N!<@<9
>1Y0!<M&@<G!<H!<F.6:627X-0+!<R+NHV.FX!2!<M&N&MWQZ?>T.R\$<G&VZ875=A!<L.M?<9L~LZ=>B1XW9308D/BBDC9X+2+M#D>P/>N%\$<J!<M+V0C!<T>T0!<T<C!<@<S-
M@TQ@<(W#<+YH!<3Y<~!<1%Z!<N!<N!<MKRC.WT!<P.DC&S!<OV@EG7T!<HJZ?<NOJ0B.N.V&V-FNOB&X0190\$<0+<B.DM613T~2~4BJC5SM.E.8@<H!<K&E+9@<CM#<I?<
90?Q6E@<T~<GRN!<WTF!<UO=<MBAW!<R~<Z~<XWA~<0PP&G!<+<#<S!<WSSWR.OJ76~<WZFWZ~<Y&Q~<6=02U4X!<M.C.55G30@<C25SM&CKNBU<~!<F8~!<F8Y!
>H5J0!<R~<57\$~<W8+!<1~<2224M53~<P~<V.VXO!<N5~<N5~<15L5&3&TDPW&D\$H4EQ!<@<9~<H!<R5~<M!<VTFP!<UDC\$!<1#~<W!<25UW&K4!<0!<P!<G5T~<1~<Q!<X!<D89~<5!<#<G0X6~<@<G!
M2I3G&3<~<J0U&N5<1NHW!<A!<5Y(7D=<8?<T!<M000A.4H4H!<@<S2~<Y!<L!<N!<P&M8+1~<R>R>ZIGC!<~<CD!<0@<?<67Q23!<3!<M!<H!<F&B&W5~<2Y=&

[illegible]

U7ZV-2M-2Y2HI-2V2EJ2KWC-GY-1>O-1(4FC6).GP7LN-MCL-MV-8/9A-SP9PL%W-D-1-GR-5Z7QV-2-(R-2<L-A5UE1%J-5EW@-MBUQ1Q1I-5-E-H-4W-TOSNEQM%
Y3ZL30GB-WKRBX85MBAL-CWI-1-MN<MK%>FVS@+MO-D-WZFSLEEF%Y-6-W8J1-RN1-2%6-MU2-1-RN<H5LQ13D-IU9T%YW-Z/H3,8<7J9S&H-5<2O-WX5X->D)=A-FW
MH00%Y=TG5-1F-414D4-B-1276-LH->IPNLIS%WFE-VX&K&KX-SOC1-WE3DVS-WP8MJOHNH3F-3K<C<X<B53>3D-HH&X0F7&T-H5-M9P00Q/JX>AD5*W0EF?
K0<-4L%KX&K>=>RDC7RQWRV5OIMU%>C<Y-MY8F6 M8%ML E(L<34-NH1=PTN/AIAK%>7P2-1W<#<Q<827?DT=@@-1-M-H-WR-M%IS6<#<
-#<@-R-A1>1H1F>T<30%P1>-Q7-PHD-VX9E M<J1F>00HV<38>H(L<1X+W00I-WMDJ(Q/P/D<Z3-PAIHSCL<U6<34%YKZ-4W<K&S&IW9C)<W/GC
MWJ9%6074=4A.#<#<=<S90?R/LCEI%>L-M-OMX-M%VWVH71Z%2%<7%<Q/L/WPPAP/Y.F%DL-1#<0<Z-BEMLA2IP2>M%EE1Q<O>D6?
03N1H-2+93MJ1%+@<S65?7PB<M<+>Q13PIPV-N\$-<OGME-MD3H<2P>=>O1G=FJX68?>2<Z<JWM-VW>Z-T&NR-1/OETV-INZRL(B2B-M>YW-78M<T-Z+RZU-
60CFL<GRS0%<#<DPT<=>R129XIO4<+<K&OBIU<K&R M-9<2<SD<0FK/ZW7%<X97<1G&H&QO F M<=>SGOV3FUEZB-NI&X&H&D&B<H I MIA&T1&83-X/OZ/IQ<@S0XN>Z1=8B<+
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GIRLZ5M\$=<EBVTNE4PDNB6X&SCHMR<F+0 Y-WA/O+6 MJHJ>+>T@6:3GZ4+G0D6<+<MHS6=1%<O/V1<2<QO.R&E(0 Y-U<HJKE7YIOEBWR5W...+WIZ-N7-MET?O?<
LOG<+>3V<1C%<#<SD3LWJ+M1R1VO-VJ3 Z-GM<F+IHGIG7XIH31<#<2B2?>P12D&A.LMUS&F6N<L-UZ8 MZK7L1L3H<H.2D4WB.KD-@09-PM57B%D.#X8Q-<
H424.CBIAP1-RC<MIFEUS&B0179L7BY>Z=<#M&F&S:XOZSO<Y%<FV&N8Z.N9G0T-CX3YIX<O-MD-72 Z1V57(SVWVHX&0/H?<@.RNHM-202C5.813L.U@G*H0SOQ=V:6XIRN
M8=72-VJW. X=E.XXINBS<+<X&H50B1YIH1BNF<7&L<+<8F1D<O-M+<1X1<1L.N#<HJZ-V-Z:V:C0817
&YPP+V-BGN>V/S3<37CMQO<C.G0&9% M22LOR42GIDJKEINB0-U.F%<WW6<HJH4&CNJ5>T MHIR&N->?FSZ?<G*E*5J1O<0#8>IH@<K&NPA<M-MH1:O2F1@?
MZRSF<1O2<QIXQ/OKB+HYHVS9L/NXV(D2-D-1?<VJO M-O-UT+P04<=>1059UQEJ\$<4O HSA<G<O <YU/F(MC-R#V M5O N->
D1-U<F<0VBR Z-D4N+X3L+VZ6<L9C.W-VL65UJOJ2.H3?M<6<=>J(OC&E&D#1<2R1<L<S&B<8509<4M3KHI-1?8HEU<H<?<F1G
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N^N^#1<2+3X+WRV#1?I<=>G(7<Y<C4B3Z1>CCT91/GV M1&O<O=>PKS1<OHT01?<COJ/A1ZQ9LMFWIHC=<MOS22B8<1-M?F2+SCW?9>01?M?2<+OSE>S0G1-U
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J<185P<+Y71J2?<Q2H<=>#<#1<J&K&R\$<MOB%000G<#<73B&FHUJ<P<=>C<@H002V7V?<+V-Y-F-HQH3<MGLQH2>ODS(C0<C Z3JW<=>Y6KJ?Z
DAXG<4CEUZO.S<=>PYYG12O.M<=>XG5OZ<F<03 JAY<@.GMJL<1O>55016<L&N<?<7-2=>S(3&H41 MNZ<U<O>OYT<D 9GH.1N8<6GWA93>C<4V RX/SHC<T1.HH?<
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JE:<5V5\$<4BV1VZ621 MK&SK/F-29<D WY%<P6JX<8AQ?7WTG<T8-%18P5TZ<SD(2RDHH=841) M(D\$3<+<HG2+
H=2N<GZUEBV-FV&8<1?8\$<U.SAX<O1J9>X<=>K&8P1=<1 MC&V1&B-1VW<+1<F<G-T5?B1.W<=>Z0&X&1&U/AC7<V.H3ANP?B-75 OM6<4XWHN-MIA01>LD6-<
9>@1H1A-M#<+1?<TK<=>@TXRH1Q?32.2949%<X-A&K<8TP<6&SPQCLEZINS<QX<X<X<I-W0H0VSVZGL T.MI<H<+<4&1QOQIB?1J-T7&S&M&INM1F=E.EVG<C<
HWSF?<U<KINK H-M?Z90<ZOG<C<5H<H VZ9M9001<@.N7<=>N4TRC<H&KM<-W7HOK<B-M#S4 GB<9S<S ZHMM1T107K9<2+8LS%<Y8=P7<C87T 9EBE
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YHVZ&ASQ3M.CV16<+<O1)MPE.HK6<ZBR4<Z&KM2>1 MJ+XK5HV+<192&O1O1V<=>1544MN.313Q<+<AZ31&3?<QHM-HRKN#7?<H1+WY351#=>PFT?>1HWDYXG.&S
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E1V1C<N.U=<W1.V16U-MRDF<121*=>S<C3ZO)<@B1*KY8P9XW<H.L-SZHG>IAUUPVNV9IH2 71=MA(OPMO>V7N%WZ1O/PAM1+MPE<#1A.XRGE.51N5VPPIN>?>(W4/
MHYK<=>A<1?<4KN?1@15TL<19PC11H(2<Y&G<FHO6YQ<R.YCJ&60A9U1H?2E M<=>1%<452T1HG2HM1FX&6<51%5Z6P1E-W?<+<M&CB9&C<O<U&R<?>1J9
M>B1N?1+H0D1A1=NSER51?R>A1Z15<1.2H?0187L2?<=>18&9?1CY9 M1P<RT1+<O2G(EB&F&F&5&81<=>6L&81HM<M<RN7Z.SP19

ABH>.MTMLD)<#CPEXHE# 6YTN R0X9NIA23=>##X"7WO>B" M#JH#XR.<V=H4THE2MR#75UO"SU2%G\$1B<KD.PJ=VXRS1=5DOXYTN>.A18(LB*Y<1N)S4FB+067
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M=6ZPC#1(M\$#P\$4RGHVAR1<\$S*JR-1B/P2Q#B.SUG<E8Y\$D#N2SIA8!M#<CK51M96/D2149BB<R5)D&J!<FR8J9D/7ZAFK62*21B8ZQW1Y71\$1QR36
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16WC+&1#>P#>.&@Y1G<100<4YJHMXN TNG#2B+1K7W)T=>B&S#Y.M#123=69W741+&3BSGJ#>1#>1#>UDB:<62\$U9R#>8L#9J7K<U-1<1>.(A2.M.K)G2;W7><ZIE%22Y16<1
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(7P2).8B>U90?<U4J#F7B<1E\$REC4L9>M9P166PCM.HW\$>HEA&S#Y+DPY0<8R ME.DIC0#>.#8K59J<H'G'S'GR&A<C\$9>?OFS#>H+1<G*FW)>AB+ISG#H73#A#H#<#*T
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D0%<C 51M9>G1Z'Y08G<YV=U8B667\$8C%57ZN)N.M3T4Y6<JR1R16M#<CX1B)B.T4.1B#B#JQ21'Z3R34'<A3S1'>2.3%>M&X2860<5>0<0RBVBM<B&K<=R5Y9THHE
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<8&Y\$ P6D(M41>(<60\$19UF2\$56BH>44HV.C08H1.5\$PQ63>0)B)1<241J0H+1'YVK<F MORYH.09HN-XE#VP41&H15H09Y1C%NIMJZ1<5C&XO>A1\$<1%<NIDB1'U'P<
M2N>#>1<15FAVZ>X+5YVS<Q\$Z\$?>ES&1#>U0 3\$S.K'2(PYM6D\$>4Y/SS<#<M T 6\$>XJX69B10H)BRAW74 JZ35=2C.KR&=>=>1DYPUZC\$IDVNA)T1<1\$<M.12FTS=H\$5145D5-
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MAL6H<1ODD(0J HJFGV.W539O>R\$1<(<87L%>Z<(R4T34.9)M'>#Y62 M6'KXKY7W<U O42N\$6R8BWN#WY6NK1FGP1J3DA-SF'V\$0<>9'N7CQC 4FCV=>X09XD5
M9E-B#<M'2V.R2#>L1\$T4Y536P4R1B2H6F.M#62A.21/9.T4.0X7EX9R 212 W\$<A1><ZQ&L0621\$R2N#>#>XDHMEF8 MEZ7.M#>

>Y1...< (Q)5SF%<A6S5W6 BEGHN+EU£+UP\$<R6ZB9KRD2:HU£+T+U0+M7&E.U2QQWCE+L.V.T4FZ\$5(0G)1(9<6<-8'06.VIO#M/£F/M9'60(£F)U£>U£.326F+K72
 Y129+<K>6W6220\$<L.VP1L.U0C0:ET+M££=AM\$YBHD9)CE£=U1(H8T0TODCB££>=5A1U1(1Q#MOSPD20>ZUHH£>TPOQKEH£
 :90K=RN.L5+9K<1GP19SOKV£F+U3U0EGXK.M1H2\$AX072>C=46N\$?E.9I24W1.BAU#FO+H#RE+H£C#YAI\$P(C0.H6RM.MK78)H\$N#<£0(C.RX2U)=5.MCJ%
 (4.O=ETVZ.T+M\$*H7L2P4.=T6(AY#74.T0B7C£L:UXC0I/PI+£C£?M£<DE9'5MY1#1#(YKIBL7£5NM.M5RC)70VC'7>4HC.D072391P06'£U6S7+I.MA\$C£
 £.40+5YHPQZ.TC6826.2/D£+WH3XD2KIWA..@9H£'I6'0.E=+M#<£0.MYT£#+TC+M#1#(UE/7£Y2%)D=£+CH.ON-Z.LJ#£LU(HTXJ2£(£0.3XV.F6H#N.NZQE(£7-2>T
 M.G£.1#£DH<6-M£+>9.G£HV7H£(H£A7)££:DEFDL.U'T9...S5W.VKSHR.QMDPR0Q.M3=1UP(V.MJFOWQ+Q.G.ZIK009£'FV£X.M.F#26)£B/F.#9'=BOBPP<1.VQ0IBM££EU03?
 9LDL£HE£2QJ6M6W\$M.Y.8%)(D-M£+M£(H£#<£0+£48'Z+P63£QP2B9)TFN£.1#£<XV3-2D/AH9£JBQZ£
 >O/7(£.82=5=R=7'2V)1(IQ-M£)H-R87TES80£TRY£<N+2W'25£P46.QUMQ...9NH\$'9XC7I2N.MS#+=N££.3Y'W=TO8'£S(£#6'T£.£82CU0£YBT£+£LP2N)BU'5'£A?£.MZ?
 #3H£2DQ1\$Y£EG5£G2W£ZC0ETI8\$N\$P12+D£>£V2(£(QJN-0.M>£57972D#£S;QV.M£6/KIC1X3H'£FMKW3+£L.GU£IQ>£ZIFN\$828.1£</M0X\$)3EYQ.BGB£&£V
 <£\$K£G6C6£7£+G3££ZC0BHAR>7%£&U£(9B8)£UD.M£G£2D8£E£I09£I05£M4£)TE8+£(£0£J£S2£D6S££B=£>QK.Z£(9KRW£ZG07.M2'N0B)0M£<GN=WNAM\$82£EZM£
 A4£F+V#£.U.N£"PP%£CQ#&£*BRIGN.M-WGB£(£3RA4%£=£.8+3ZSFDDWQ7S&KUHZ.TC'£F££G£:8@7°C.M£.7LSBR.M2JD£\$£.J£1H£>>ZY\$£?ZIP0OR.O£OLK\$£FFF.SAZ-£#03
 M6£6LAD>£L.MM#R£4£FKKDD<0£L£6£M6BX9£1V4U£9'£3£A52C(SXV£4P6W9HW£X+KLS.M5£IPGMYT£JCB£CFN£C£0£)=£#Q/ZXZ?A£>A0£.£.W77GN£+£.G2H£42£
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been prepared in accordance with accounting principles generally accepted in the United States of America ("GAAP") and include all adjustments necessary for the fair presentation of the Company's financial position for the periods presented.

The consolidated financial statements include the balances of the Company and its subsidiaries. All intercompany transactions and balances are eliminated in consolidation.

Unaudited Interim Financial Information

The accompanying unaudited condensed consolidated financial statements as of June 30, 2024 and for the three and six months ended June 30, 2024 and 2023, have been prepared in accordance with GAAP for interim financial statements and pursuant to Form 10-Q and Article 10 of Regulation S-X. In the opinion of management, all adjustments (consisting of normal recurring accruals) have been made that are considered necessary for a fair statement of the financial position of the Company as of June 30, 2024, the results of operations for the three and six months ended June 30, 2024 and 2023 and the cash flows for the six months ended June 30, 2024 and 2023. The condensed consolidated balance sheet as of December 31, 2023 has been derived from the Company's audited consolidated financial statements. Operating results for the three and six months ended June 30, 2024 are not necessarily indicative of the results that may be expected for the year ending December 31, 2024.

Certain information and disclosures normally included in the notes to annual financial statements prepared in accordance with GAAP have been omitted from these interim financial statements pursuant to the rules and regulations of the Securities and Exchange Commission (the "SEC"). Accordingly, these unaudited condensed consolidated financial statements should be read in conjunction with the financial statements for the fiscal year ended December 31, 2023, which are included in the Company's 10-K filed with the SEC on February 29, 2024.

Liquidity

As of June 30, 2024, the Company has an accumulated deficit of approximately \$820.1 million and net cash used in operating activities was approximately \$53.0 million for the six months ended June 30, 2024. Management expects to continue to incur operating losses and negative cash flows from operations for the foreseeable future.

As of June 30, 2024, the Company had cash, cash equivalents, and marketable securities of \$577.2 million. The Company believes that its existing cash, cash equivalents, and marketable securities will be sufficient to meet its cash commitments for at least the next 12 months after the date that these consolidated financial statements are issued. The Company's research and development and preparation for potentially becoming a commercial stage organization can be costly, and the timing and outcomes are uncertain. The assumptions upon which the Company has based its estimates are routinely evaluated and may be subject to change. The actual amount of the Company's expenditures will vary depending upon a number of factors including but not limited to the progress of the Company's research and development activities and potential regulatory approval of its product candidates and the level of financial resources available.

Significant Risks and Uncertainties

The Company's operations are subject to a number of factors that can affect its operating results and financial condition. Such factors include, but are not limited to: the results of research and development, clinical testing and trial activities of the Company's products, the Company's ability to obtain regulatory approval to market its products, competition from products manufactured and sold or being developed by other companies, the price of, and demand for, Company's products, the Company's ability to negotiate favorable licensing or other manufacturing and marketing agreements for its products, and the Company's ability to raise capital.

The Company currently has no commercially approved products and there can be no assurance that the Company's research and development will be successfully commercialized. Developing and commercializing a product requires significant time and capital and is subject to regulatory review and approval as well as competition from other biotechnology and pharmaceutical companies. The Company operates in an environment of rapid change and is dependent upon the continued services of its employees and vendors and obtaining and protecting intellectual property.

Use of Estimates

The preparation of financial statements in conformity with GAAP requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and the disclosure of contingent assets and liabilities at the date of the financial statements and the reported amount of revenue and expenses during the year. Accordingly, actual results could differ from those estimates and those differences could be significant. Significant estimates and assumptions reflected in the accompanying consolidated financial statements include, but are not limited to: warrant liabilities, leases, stock options granted and depreciation expense.

Functional Currency and Foreign Currency Translations

The functional currency of the Company's subsidiaries incorporated in the People's Republic of China ("PRC") is Renminbi ("RMB"). The functional currency of the Company and its subsidiaries incorporated outside the PRC is USD. Transactions denominated in currencies other than the functional currencies are re-measured into the functional currency of the entity at the exchange rates prevailing on the transaction dates. Foreign currency denominated financial assets and liabilities are re-measured at the balance sheet date exchange rate. The resulting exchange differences are included in the net loss of the statements of operations and comprehensive loss. Assets and liabilities of the Company with functional currency other than USD are translated into USD at fiscal year-end exchange rates. Equity amounts are translated at historical exchange rates. Income and expense items are translated at average exchange rates during the fiscal year. Translation adjustments arising from these are reported as foreign currency translation adjustments and are shown as a component of other comprehensive income (loss).

Cash and Cash Equivalents

Cash equivalents include short-term, highly liquid instruments, consisting of money market accounts, a money market mutual fund and short-term investments with maturities from the date of purchase of 90 days or less. The majority of cash and cash equivalents are maintained with major financial institutions in North America. Deposits with these financial institutions may exceed the amount of insurance provided on such deposits. These deposits may be redeemed upon demand which reduces counterparty performance risk.

Marketable Securities

Debt securities have been classified as available-for-sale which may be sold before maturity or are not classified as held to maturity or trading. Marketable debt securities classified as available-for-sale are carried at fair value with unrealized gains or losses reported in other comprehensive income (loss). Exchange traded funds are equity securities, which are reported as marketable securities with readily determinable fair values, are also carried at fair value with unrealized gains and losses included in other income (expense), net. Realized gains and losses on both debt and equity securities are included in other income (expense), net.

For securities in an unrealized loss position, management considers the extent and duration of the unrealized loss, and the financial condition and near-term prospects of the issuer. Management also assesses whether it intends to sell, or it is more likely than not that it will be required to sell, a security in an unrealized loss position before recovery of its amortized cost basis. If management determines there is any other than temporary impairment, the entire difference between amortized cost and fair value is recognized as impairment through earnings.

Debt securities

The Company is exposed to credit losses primarily through its available-for-sale investments. The Company assesses whether its available-for-sale investments are impaired at each reporting period. Unrealized losses or impairments resulting from the amortized cost basis of any available-for-sale debt security exceeding its fair value are evaluated for identification of credit losses and non-credit related losses. Any credit losses are charged to earnings against the allowance for credit losses of the debt security, limited to the difference between the fair value and the amortized cost basis of the debt security. Any difference between the fair value of the debt security and the amortized cost basis, less the allowance for expected credit losses, are reported in other comprehensive income (loss). Expected cash inflows due to improvements in credit are recognized through a reversal of the allowance for expected credit losses subject to the total allowance previously recognized. The Company's expected loss allowance methodology for the debt securities includes reviewing the extent of the unrealized loss, the size, term, geographical location, and industry of the issuer, the issuers' credit ratings and any changes in those ratings, as well as reviewing current and future economic market conditions and the issuers' current status and financial condition. As of June 30, 2024, the Company has not recognized an allowance for expected credit losses related to available-for-sale investments as the Company has not identified any unrealized losses for these investments attributable to credit factors.

Interest income

Interest income includes amortization and accretion of purchase premiums and discounts. Premiums and discounts on debt securities are amortized on the effective-interest method. Gains and losses on sales are recorded on the settlement date and determined using the specific identification method.

Concentration of Risk

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wrap;font-family:'Calibri',sans-serif;min-width:fit-content;"></p></td></tr></table></div><div style="font-size:12pt;font-family:Times New Roman;"><p style="font-size:10pt;margin-top:12pt;margin-bottom:0;text-align:justify;">Property and Equipment</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Property and equipment are stated at cost less accumulated depreciation. Property and equipment are depreciated using the straight-line method over the estimated useful lives of the related assets of generally five yearsfor computers andseven years</p></div><div style="font-size:10pt;font-family:Times New Roman;"><p style="font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Impairment of Long-Lived Assets</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The Company reviews long-lived assets for impairment whenever events or changes in circumstances indicate that the carrying amount of an asset may not be recoverable and an impairment assessment may be performed on the recoverability of the carrying amounts. If an impairment occurs, the loss is measured by comparing the fair value of the asset to its carrying amount.</p></div><div style="font-size:10pt;font-family:Times New Roman;"><p style="font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Warrants</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The Company accounts for warrants as either equity-classified or liability-classified instruments based on an assessment of the warrant's specific terms and applicable authoritative guidance in Financial Accounting Standards Board ("FASB") Accounting Standards Codification ("ASC") 480, Distinguishing Liabilities from Equity ("ASC 480") and ASC 815, Derivatives and Hedging ("ASC 815"). The assessment considers whether the warrants are freestanding financial instruments pursuant to ASC 480, meet the definition of a liability pursuant to ASC 480, and whether the warrants meet all of the requirements for equity classification under ASC 815, including whether the warrants are indexed to the Company's own ordinary shares, among other conditions for equity classification. This assessment, which requires the use of professional judgment, is conducted at the time of warrant issuance and as of each subsequent quarterly period end date while the warrants are outstanding.</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">For issued or modified warrants that meet all of the criteria for equity classification, the warrants are required to be recorded as a component of additional paid-in capital at the time of issuance. For issued or modified warrants that do not meet all the criteria for equity classification, the warrants are required to be recorded as a liability at their fair value on the date of issuance and each balance sheet date thereafter. Changes in the estimated fair value of the warrants are recognized as a non-cash gain or loss on the statements of operations and comprehensive loss.</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Following the Merger, there were 5,787,472 warrants to purchase Common Stock outstanding, consisting of 4,791,639 Public Warrants,162,500 Private Placement Warrants and833,333 Forward Purchase Warrants. Each whole warrant entitles the registered holder to purchase one share of our Class A Common Stock at a price of \$11.50 per share. Pursuant to the warrant agreement, a warrant holder may exercise its warrants only for a whole number of shares of our Class A Common Stock.</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">As part of the consideration for AnHeart, the Company issued warrants collectively exercisable for approximately 2,893,731 shares of the Company's Class A Common Stock at an exercise price of \$11.50 per share (the "Consideration Warrants"). The fair value of the Consideration Warrants was determined to approximate the public warrant trading price as of closing of the Acquisition, yielding a fair value of approximately \$0.26 per share. The Consideration Warrants are restricted with respect to the exercise and transfer thereof until receipt of such stockholder approval and otherwise have terms identical to those of the Company's outstanding Public Warrants. Accordingly, the Company classified the Consideration Warrants as liabilities, consistent with the classification of the Company's Public Warrants.</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The Company evaluated Public Warrants, Private Placement Warrants, Consideration Warrants, and Forward Purchase Warrants (the "Warrants") under ASC 815-40, Derivatives and Hedging—Contracts in Entity's Own Equity, and concluded that they do not meet the criteria to be classified in stockholders' equity. Specifically, the settlement value of the Warrants is dependent, in part, on the holder of the Warrants at the time of settlement. Because the holder of an instrument is not an input into the pricing of a fixed-for-fixed option on our Common Stock, the Warrants fail the indexation guidance in ASC 815-40, which would preclude classification in stockholders' equity. Additionally,the exercise of the Warrants may be settled in cash upon the occurrence of a tender offer or exchange that involves more than 50% of the outstanding shares of the Company's Common Stock.Because not all of the Company's stockholders need to participate in such tender offer or exchange to trigger the potential cash settlement and the Company does not control the occurrence of such an event, the Company concluded that the Warrants do not meet the conditions to be classified in equity. Since the Warrants meet the definition of a derivative under ASC 815, the Company recorded these Warrants as liabilities on the balance sheet at fair value upon the closing of the Merger, with subsequent changes in their respective fair values recognized in the consolidated statement of operations and comprehensive loss at each reporting date. The fair value of Consideration, Public, and Forward Purchase Warrants was determined using the closing price of the publicly traded warrants on the NYSE market. The fair value of the Private Placement Warrants was estimated using a Black-Scholes option pricing model(see Note 4)</p></div><div style="font-size:12pt;font-family:Times New Roman;"><p style="font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Revenue Recognition</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The Company applies ASC Topic 606, "Revenue from Contracts with Customers" ("ASC 606") to account for its revenue transactions. Under ASC 606, an entity recognizes revenue when its customer obtains control of promised goods or services, in an amount that reflects the consideration that the entity expects to receive in exchange for those goods or services. To determine revenue recognition for arrangements that an entity determines are within the scope of ASC 606, the entity performs the five-step model. The Company only applies the five-step model to contracts when it is probable that the entity will collect the consideration to which it is entitled in exchange for the goods or services it transfers to the customer. Once a contract is determined to be within the scope of ASC</p><p style="font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">606at contract inception, the Company reviews the contract to determine which performance obligations it must deliver and which of these performance obligations are distinct. The Company recognizes as revenue the amount of the transaction price that is allocated to each performance obligation when that performance obligation is satisfied or as it is satisfied. Accounts receivable represent amounts invoiced and revenues recognized prior to invoicing when the Company has satisfied its performance obligation and has the unconditional right to payment.</p></div><div style="font-size:12pt;font-family:Times New Roman;"><p style="font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Collaborative Arrangement</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">In November 2018, the FASB issued Topic 808, "Collaborative Arrangements" ("ASC 808"). Clarifying the Interaction between ASC 808 and ASC 606. This update clarifies that certain transactions between participants in a collaborative arrangement should be accounted for under ASC 606 when the counterparty is a customer and precludes an entity from presenting consideration from a transaction in a collaborative arrangement as revenue from contracts with customers if the counterparty is not a customer for that transaction. The Company adopted this standard for all periods presented. The Company enters into collaborative arrangements for the research and development, manufacture and/or commercialization of drug products and drug candidates. The Company assessed and determined that none of the collaboration agreements entered into during the periods presented were within the scope of ASC 808, as all of the agreements did not involve active participation by both parties in a joint research activity, and therefore did not qualify as collaborative arrangements under ASC 808. The Company has determined that all the elements of the above collaborations are reflective of a vendor-customer relationship and therefore within the scope of ASC 606. In determining the appropriate amount of revenue to be recognized as it fulfills its obligations under each of its agreements, the Company performs the five-step model under ASC 606 noted above. The Company recognizes revenue to depict the transfer of control of promised goods or services to customers in an amount that reflects the consideration to which the entity expects to receive in exchange for those goods or services. The Company's collaborative arrangements may contain more than one unit of account, or performance obligation, including grants of licenses to intellectual property rights, agreements to provide research and development services and other deliverables. The collaborative arrangements do not include a right of return for any deliverable. As part of the accounting for these arrangements, the Company must develop assumptions that require judgment to determine the stand-alone selling price for each performance obligation identified in the contract. In developing the stand-alone selling price for a performance obligation, the Company considers competitor pricing for a similar or identical product, market awareness of and perception of the product, expected product life and current market trends. In general, the consideration allocated to each performance obligation is recognized when the respective obligation is satisfied either by delivering a good or providing a service. Limited to the consideration that is not constrained. Non-refundable payments received before all of the relevant criteria for revenue recognition are satisfied are recorded as contract liabilities.</p><p style="font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Licenses of intellectual property</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">If a license to the Company's intellectual property is determined to be distinct from the other promises or performance obligations identified in the contract, the Company recognizes revenue from the portion of the transaction price allocated to the license at a point in time, when the license is

transferred to the customer and the customer is able to benefit from the license.

Research and development services

The portion of a transaction price allocated to research and development services performance obligations is deferred and recognized as revenue over time as delivery or performance of such services occurs based on the use of an input method.

Milestones payments

Customer options

The Company's collaboration agreements include development and regulatory milestones. The Company evaluates whether the milestones are considered probable of being reached and estimates the amount to be included in the transaction price using the most likely amount method. The Company evaluates factors such as the scientific, clinical, regulatory, commercial, and other risks that must be overcome to achieve the particular milestone in making this assessment. If it is probable that a significant revenue reversal would not occur, the associated milestone value is included in the transaction price. Milestone payments that are not within the Company's control or the licensee's control, such as regulatory approvals, are not considered probable of being achieved until those approvals are received. At the end of each reporting period, the Company re-evaluates the probability of achievement of such milestones and any related

Royalties

For sales-based royalties, including milestone payments based on the level of sales, the Company determines whether the sole or predominant item to which the royalties relate is a license. When the license is the sole or predominant item to which the sales-based royalty relates, the Company recognizes revenue at the later of: (i) when the related sales occur, or (ii) when the performance obligation to which some or all of the royalty has been allocated has been satisfied (or partially satisfied). To date, the Company has not recognized any sales-based royalty revenue resulting from the Company's collaboration agreements.

The Company receives payments from its customers based on billing terms established in the contract. Up-front payments and fees are recorded as contract liabilities (e.g., deferred revenue) upon receipt or when due until the Company performs its obligations under the arrangement.

Costs to Fulfill a Contract with a Customer

The Company has not capitalized any costs for the three and six months ended June 30, 2024 and 2023. The Company is required to capitalize costs incurred to fulfill customer contracts. These costs are required to be amortized to expense on a systemic basis that is consistent with the transfer to the customer of the goods or services to which the asset relates, compared to previously being expensed as incurred. Elements of the costs primarily include (i) payroll and other related costs of personnel related directly to the contract activities; (ii) costs related to pre-clinical testing and clinical trials such as payments to contract research organizations ("CROs") and contract manufacturing organizations ("CMOs"), investigators, and clinical trial sites that conduct the contract activities; (iv) costs to develop the product candidates, including raw materials and supplies, product testing, depreciation, and facility-related expenses; and (v) other research and development costs.

Leases

Operating lease ROU assets and operating lease liabilities are recognized based on the present value of the future minimum lease payments over the lease term at the commencement date. ROU assets also include any initial direct costs incurred and any lease payments made on or before the lease commencement date, less lease incentives received. The Company uses its incremental borrowing rate based on the information available at the commencement date in determining the lease liabilities as the Company's leases generally do not provide an implicit rate. The incremental borrowing rate is reevaluated upon a lease modification. The operating lease ROU asset also includes any initial direct costs and prepaid lease payments made less any lease incentives. The Company considered information available at the adoption date of Topic 842 to determine the incremental borrowing rate for leases in existence as of this date. The incremental borrowing rate used was the weighted average rate between secured and unsecured lending arrangements. Lease terms may include options to extend or terminate the lease when the Company is reasonably certain that the option will be exercised. Variable payments included in the lease agreement are expensed as incurred. Lease expense is recognized on a straight-line basis over the lease term.

Segment Information

The Company manages its operations as a single segment for the purposes of assessing performance and making operating decisions. The Company's operations are focused on oncology development activities.

Research and Development Costs

Research and development activities are expensed as incurred. These costs include fees paid to consultants, vendors and various entities that perform certain research and testing on behalf of the Company.

Stock-based Compensation

The Company recognizes compensation cost for grants of employee stock options using a fair value measurement method, that is recognized in operating results as compensation expense based on the fair value over the requisite service period of the awards. Forfeitures are recorded as they occur instead of estimating forfeitures that are expected to occur.

Black-Scholes option-pricing model

The Company determines the fair value of stock-based awards that are based only on a service condition using the Black-Scholes option-pricing model which uses both historical and current market data to estimate fair value. The method incorporates various assumptions such as the risk-free interest rate, volatility, dividend yield, and expected life of the options.

Income Taxes

The Company determines the fair value of stock-based awards that are based on both a service condition and achievement of the first to occur of a market or performance condition using a Monte Carlo simulation.

Income Taxes

The Company accounts for income taxes using the asset and liability method, which requires the recognition of deferred tax assets and liabilities for the expected future tax consequences of events that have been recognized in the consolidated financial statements or in the Company's tax returns. The difference between the financial statement and tax basis of assets and liabilities is determined annually. Deferred income tax assets and liabilities are computed for those differences that have future tax consequences using the currently enacted

tax laws and rates that apply to the years in which they are expected to affect taxable income. The Company assesses the likelihood that its deferred tax assets will be recovered from future taxable income and, to the extent it believes, based upon the weight of available evidence, that it is more likely than not that all or a portion of the deferred tax assets will not be realized, a valuation allowance is established through a charge to income tax expense.

10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The Company uses a recognition threshold and measurement attribute for the financial statement recognition and measurement of a tax position taken, or expected to be taken, in a tax return. The provision for income taxes includes the effects of any resulting tax reserves, or unrecognized tax benefits, that are considered appropriate. The Company's policy is to record interest and penalties related to income taxes as part of the tax provision. Returns for tax years beginning with those filed for the period ended December 31, 2018 are open to federal and state tax examination.

10pt;font-family:Times New Roman;"><p style="font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Recent Accounting Pronouncements</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">In November 2023, the FASB issued Topic 280 "Improvements to Reportable Segment Disclosures" which expands annual and interim disclosure requirements for reportable segments, primarily through enhanced disclosures about significant segment expenses. This guidance is effective for our annual periods beginning January 1, 2024, and for interim periods beginning January 1, 2025, with early adoption permitted. We are currently evaluating the potential effect that the updated standard will have on our financial statement disclosures.

10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">In December 2023, the FASB issued Topics 740 "Improvements to Income Tax Disclosures" to expand the disclosure requirements for income taxes, specifically related to the rate reconciliation and income taxes paid. This guidance is effective for our annual periods beginning January 1, 2025, with early adoption permitted. We are currently evaluating the potential effect that the updated standard will have on our financial statement disclosures.

10pt;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Basis of Presentation</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The accompanying consolidated financial statements have been prepared in accordance with accounting principles generally accepted in the United States of America ("GAAP") and include all adjustments necessary for the fair presentation of the Company's financial position for the periods presented.

10pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Principles of Consolidation</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The consolidated financial statements include the balances of the Company and its subsidiaries. All intercompany transactions and balances are eliminated in consolidation.

10pt;font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Unaudited Interim Financial Information</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The accompanying unaudited condensed consolidated financial statements as of June 30, 2024 and for the three and six months ended June 30, 2024 and 2023, have been prepared in accordance with GAAP for interim financial statements and pursuant to Form 10-Q and Article 10 of Regulation S-X. In the opinion of management, all adjustments (consisting of normal recurring accruals) have been made that are considered necessary for a fair statement of the financial position of the Company as of June 30, 2024, the results of operations for the three and six months ended June 30, 2024 and 2023 and the cash flows for the six months ended June 30, 2024 and 2023. The condensed consolidated balance sheet as of December 31, 2023 has been derived from the Company's audited consolidated financial statements. Operating results for the three and six months ended June 30, 2024 are not necessarily indicative of the results that may be expected for the year ending December 31, 2024.

10pt;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Certain information and disclosures normally included in the notes to annual financial statements prepared in accordance with GAAP have been omitted from these interim financial statements pursuant to the rules and regulations of the Securities and Exchange Commission (the "SEC"). Accordingly, these unaudited condensed consolidated financial statements should be read in conjunction with the financial statements for the fiscal year ended December 31, 2023, which are included in the Company's 10-K filed with the SEC on February 29, 2024.

10pt;font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Liquidity</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">As of June 30, 2024, the Company has an accumulated deficit of approximately

10pt;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">ion and net cash used in operating activities was approximately

10pt;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">53.0

10pt;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">million. The Company believes that its existing cash, cash equivalents, and marketable securities will be sufficient to meet its cash commitments for at least the next 12 months after the date that these consolidated financial statements are issued. The Company's research and development and preparation for potentially becoming a commercial stage organization can be costly, and the timing and outcomes are uncertain. The assumptions upon which the Company has based its estimates are routinely evaluated and may be subject to change. The actual amount of the Company's expenditures will vary depending upon a number of factors including but not limited to the progress of the Company's research and development activities and potential regulatory approval of its product candidates and the level of financial resources available.

10pt;font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Significant Risks and Uncertainties</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The Company's operations are subject to a number of factors that can affect its operating results and financial condition. Such factors include, but are not limited to: the results of research and development, clinical testing and trial activities of the Company's products, the Company's ability to obtain regulatory approval to market its products, competition from products manufactured and sold or being developed by other companies, the price of, and demand for, Company's products, the Company's ability to negotiate favorable licensing or other manufacturing and marketing agreements for its products, and the Company's ability to raise capital.

10pt;font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The Company currently has no commercially approved products and there can be no assurance that the Company's research and development will be successfully commercialized. Developing and commercializing a product requires significant time and capital and is subject to regulatory review and approval as well as competition from other biotechnology and pharmaceutical companies. The Company operates in an environment of rapid change and is dependent upon the continued services of its employees and vendors and obtaining and protecting intellectual property.

10pt;font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Use of Estimates</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The preparation of financial statements in conformity with GAAP requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and the disclosure of contingent assets and liabilities at the date of the financial statements and the reported amount of revenue and expenses during the year. Accordingly, actual results could differ from those estimates and those differences could be significant. Significant estimates and assumptions reflected in the accompanying consolidated financial statements include, but are not limited to: warrant liabilities, leases, stock options granted and depreciation expense.

10pt;font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Functional Currency and Foreign Currency Translations</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The functional currency of the Company and its subsidiaries incorporated outside the PRC is USD. Transactions denominated in currencies other than the functional currencies are re-measured into the functional currency of the entity at the exchange rates prevailing on the transaction dates. Foreign currency denominated financial assets and liabilities are re-measured at the balance sheet date exchange rate. The resulting exchange differences are included in the net loss of the statements of operations and comprehensive loss. Assets and liabilities of the Company with functional currency other than USD are translated into USD at fiscal year-end exchange rates. Equity amounts are translated at historical exchange rates. Income and expense items are translated at average exchange rates during the fiscal year. Translation adjustments arising from these are reported as foreign currency translation adjustments and are shown as a component of other comprehensive income (loss).

10pt;font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Cash and Cash Equivalents</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Cash equivalents include short-term, highly liquid instruments, consisting of money market accounts, a money market mutual fund and short-term investments with maturities from the date of purchase of 90 days or less. The majority of cash and cash equivalents are maintained with major financial institutions in North America. Deposits with these financial institutions may exceed the amount of insurance provided on such deposits. These deposits may be redeemed upon demand which reduces counterparty performance risk.

10pt;font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Marketable Securities</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Debt securities have been classified as available-for-sale which may be sold before maturity or are not classified as held-to-maturity or trading. Marketable debt securities classified as available-for-sale are carried at fair value with unrealized gains or losses reported in other comprehensive income (loss). Exchange traded funds are equity securities, which are reported as marketable securities with readily determinable fair values, are also carried at fair value with unrealized gains and losses included in other income (expense), net. Realized gains and losses on both debt and equity securities are included in other income (expense), net.

10pt;font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">For securities in an unrealized loss position, management considers the extent and duration of the unrealized loss, and the financial condition and near-term prospects of the issuer. Management also assesses whether it intends to sell, or it is more likely than not that it will be required to sell, a security in an unrealized loss position before recovery of its amortized cost basis. If management

10pt;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">etermines there is any

[illegible]

[illegible]

[illegible]

research and development services and other deliverables. The collaborative arrangements do not include a right of return for any deliverable. As part of the accounting for these arrangements, the Company must develop assumptions that require judgment to determine the stand-alone selling price for each performance obligation identified in the contract. In developing the stand-alone selling price for a performance obligation, the Company considers competitor pricing for a similar or identical product, market awareness of and perception of the product, expected product life and current market trends. In general, the consideration allocated to each performance obligation is recognized when the respective obligation is satisfied either by delivering a good or providing a service, limited to the consideration that is not constrained. Non-refundable payments received before all of the relevant criteria for revenue recognition are satisfied are recorded as contract liabilities.

A customer's right to choose, at its discretion, to make a payment for additional goods or services is generally considered an option. If the Company is not presently obligated to provide and does not have a right to consideration for delivering additional goods or services, the item is considered an option. The Company evaluates the customer options for material rights, such as the ability to acquire additional goods or services for free or a discount. Optional future services that reflect their standalone selling prices do not provide the customer with a material right and, therefore, are not considered performance obligations and are accounted for as separate contracts. The optional future services do not include a material right to be accounted for as performance obligations.

A customer's right to choose, at its discretion, to make a payment for additional goods or services is generally considered an option. If the Company is not presently obligated to provide and does not have a right to consideration for delivering additional goods or services, the item is considered an option. The Company evaluates the customer options for material rights, such as the ability to acquire additional goods or services for free or a discount. Optional future services that reflect their standalone selling prices do not provide the customer with a material right and, therefore, are not considered performance obligations and are accounted for as separate contracts. The optional future services do not include a material right to be accounted for as performance obligations.

Milestones payments

The Company's collaboration agreements include development and regulatory milestones. The Company evaluates whether the milestones are considered probable of being reached and estimates the amount to be included in the transaction price using the most likely amount method. The Company evaluates factors such as the scientific, clinical, regulatory, commercial, and other risks that must be overcome to achieve the particular milestone in making this assessment. If it is probable that a significant revenue reversal would not occur, the associated milestone value is included in the transaction price. Milestone payments that are not within the Company's control or the licensee's control, such as regulatory approvals, are not considered probable of being achieved until those approvals are received. At the end of each reporting period, the Company re-evaluates the probability of achievement of such milestones and any related

Royalties

For sales-based royalties, including milestone payments based on the level of sales, the Company determines whether the sole or predominant item to which the royalties relate is a license. When the license is the sole or predominant item to which the sales-based royalty relates, the Company recognizes revenue at the later of: (i) when the related sales occur; or (ii) when the performance obligation to which some or all of the royalty has been allocated has been satisfied (or partially satisfied). To date, the Company has not recognized any sales-based royalty revenue resulting from the Company's collaboration agreements.

Costs to Fulfill a Contract with a Customer

The Company receives payments from its customers based on billing terms established in the contract. Up-front payments and fees are recorded as contract liabilities (e.g., deferred revenue) upon receipt or when due until the Company performs its obligations under the arrangement.

Leases

The Company currently does not have any finance leases.

Operating lease ROU assets and operating lease liabilities are recognized based on the present value of the future minimum lease payments over the lease term at the commencement date. ROU assets also include any initial direct costs incurred and any lease payments made on or before the lease commencement date, less lease incentives received. The Company uses its incremental borrowing rate based on the information available at the commencement date in determining the lease liabilities as the Company's leases generally do not provide an implicit rate. The incremental borrowing rate is reevaluated upon a lease modification. The operating lease ROU asset also includes any initial direct costs and prepaid lease payments made less any lease incentives. The Company considered information available at the adoption date of Topic 842 to determine the incremental borrowing rate for leases in existence as of this date. The incremental borrowing rate used was the weighted average rate between secured and unsecured lending arrangements. Lease terms may include options to extend or terminate the lease when the Company is reasonably certain that the option will be exercised. Variable payments included in the lease agreement are expensed as incurred. Lease expense is recognized on a straight-line basis over the lease term.

Segment Information

The Company manages its operations as a single segment for the purposes of assessing performance and making operating decisions. The Company's operations are focused on oncology development activities.

Research and Development Costs

Costs incurred in connection with research and development activities are expensed as incurred. These costs include fees paid to consultants, vendors and various entities that perform certain research and testing on behalf of the Company.

Stock-based Compensation

The Company elected to apply each of the practical expedients described in ASC Topic 842-10-65-1(f) which allow companies not to reassess: (i) whether any expired or existing agreements contain leases, (ii) the classification of any expired or existing leases, and (iii) the capitalization of initial direct costs for any existing leases. The Company also elected to apply the short-term lease measurement and recognition exemption in which ROU assets and lease liabilities are not recognized for short-term operating leases. A short-term operating lease is a lease that, at the commencement date, has a lease term of 12 months or less and does not include an option to purchase the underlying asset that the lessee is reasonably certain to exercise. The Company also elected not to separate non-lease components from lease components and instead to account for each separate lease component and the non-lease components associated with that lease component as a single lease component.

Forfeitures

Forfeitures are recorded as they occur instead of estimating forfeitures that are expected to occur.

Costs incurred in connection with research and development activities are expensed as incurred. These costs include fees paid to consultants, vendors and various entities that perform certain research and testing on behalf of the Company.

The Company recognizes compensation cost for grants of employee stock options using a fair-value measurement method, that is recognized in operating results as compensation expense based on the fair value over the requisite service period of the awards. Forfeitures are recorded as they occur instead of estimating forfeitures that are expected to occur.

The Company determines the fair value of stock-based awards that are based only on a service condition using the Black-Scholes option-pricing model which uses both historical and current market data to estimate fair value. The method incorporates various assumptions such as the risk-free interest rate, volatility, dividend yield, and expected life of the options.

New Roman;margin-bottom:0;text-align:justify;">The Company determines the fair value of stock-based awards that are based on both a service condition and achievement of the first to occur of a market or performance condition using a Monte Carlo simulation.</p><p style="font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Income Taxes</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The Company accounts for income taxes using the asset and liability method, which requires the recognition of deferred tax assets and liabilities for the expected future tax consequences of events that have been recognized in the consolidated financial statements or in the Company's tax returns. The difference between the financial statement and tax basis of assets and liabilities is determined annually. Deferred income tax assets and liabilities are computed for those differences that have future tax consequences using the currently enacted tax laws and rates that apply to the years in which they are expected to affect taxable income. The Company assesses the likelihood that its deferred tax assets will be recovered from future taxable income and, to the extent it believes, based upon the weight of available evidence, that it is more likely than not that all or a portion of the deferred tax assets will not be realized, a valuation allowance is established through a charge to income tax expense.</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Recent Accounting Pronouncements</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">In November 2023, the FASB issued Topic 280 "Improvements to Reportable Segment Disclosures" which expands annual and interim disclosure requirements for reportable segments, primarily through enhanced disclosures about significant segment expenses. This guidance is effective for our annual periods beginning January 1, 2024, and for interim periods beginning January 1, 2025, with early adoption permitted. We are currently evaluating the potential effect that the updated standard will have on our financial statement disclosures.</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">In December 2023, the FASB issued Topics 740 "Improvements to Income Tax Disclosures" to expand the disclosure requirements for income taxes, specifically related to the rate reconciliation and income taxes paid. This guidance is effective for our annual periods beginning January 1, 2025, with early adoption permitted. We are currently evaluating the potential effect that the updated standard will have on our financial statement disclosures.</p><p style="font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:left;">3. Acquisition</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">On the Acquisition Date, the Company completed the Acquisition of AnHeart. Pursuant to the terms of the AnHeart Merger Agreement, at the effective time of the First Merger (as defined in the Merger Agreement) (the "First Effective Time"), the Company issued to AnHeart securityholders (i) approximately 27,646,255 shares of Class A Common Stock, (ii) 851,202 shares of Series A Non-Voting Convertible Preferred Stock of the Company, and (iii) the Consideration Warrants. The shares of our Series A Non-Voting Convertible Preferred Stock are automatically convertible into an aggregate of approximately 85,120,200 shares of Class A Common Stock upon the approval of such conversion by the Company's stockholders in accordance with the rules of the New York Stock Exchange. Since the preferred stock is not mandatorily redeemable and the exception for a deemed liquidation event was not met, all of the outstanding convertible preferred stock is classified as mezzanine equity in the balance sheet. The Consideration Warrants are restricted with respect to the exercise and transfer thereof until receipt of such stockholder approval and otherwise have terms identical to those of the Company's outstanding Public Warrants.</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">At the First Effective Time, (i) each option to purchase shares of Ordinary Shares (as defined in the Merger Agreement) of AnHeart (an "AnHeart Option") held by a Continuing Company Service Provider (as defined in the Merger Agreement), whether or not vested, was assumed and converted into an option to purchase Class A Common Stock (each such option, an "Assumed Option"), and (ii) each restricted stock unit reflecting the right to receive Ordinary Shares of AnHeart (an "AnHeart RSU") held by a Continuing Company Service Provider (as defined in the Merger Agreement) was assumed and became a restricted stock unit with respect to a number of shares of Class A Common Stock (each such restricted stock unit, an "Assumed RSU"). Each Assumed Option and Assumed RSU is subject to the same terms and conditions (including vesting and exercise schedule) as were applicable to the corresponding AnHeart Option or AnHeart RSU immediately prior to the First Effective Time, subject to limited exceptions set forth in the Merger Agreement. Any other AnHeart Option and AnHeart RSU that remained unexercised and outstanding as of immediately prior to the First Effective Time was canceled without payment. From and after the Effective Time:</p><div class="item-list-element-wrapper" style="margin-left:6.667%;display:flex;margin-top:6pt;justify-content:flex-start;align-items:baseline;margin-bottom:0;min-width:6.667%;text-align:justify;"><div style="width:100%;display:inline;">the number of shares of Class A Common Stock subject to each Assumed Option was determined by multiplying (A) the number of Ordinary Shares of AnHeart that are subject to such AnHeart Option immediately prior to the First Effective Time, by (B) the Equity Award Exchange Ratio (as defined in the Merger Agreement) and rounding down to the nearest whole number of shares;</div></div><div class="item-list-element-wrapper" style="margin-left:6.667%;display:flex;margin-top:6pt;justify-content:flex-start;align-items:baseline;margin-bottom:0;min-width:6.667%;text-align:justify;"><div style="width:100%;display:inline;">the per-share exercise price for Class A Common Stock issuable upon exercise of each Assumed Option was determined by dividing (A) the per-share exercise price of each AnHeart Option, as in effect immediately prior to the First Effective Time, by (B) the Equity Award Exchange Ratio and rounding up to the nearest whole cent; and</div></div><div class="item-list-element-wrapper" style="margin-left:6.667%;display:flex;margin-top:6pt;justify-content:flex-start;align-items:baseline;margin-bottom:0;min-width:6.667%;text-align:justify;"><div style="width:100%;display:inline;">the number of shares of Class A Common Stock subject to each Assumed RSU was determined by multiplying (A) the number of Ordinary Shares of AnHeart that are subject to such AnHeart RSU immediately prior to the First Effective Time, by (B) 1.7551 and rounding down to the nearest whole number of shares.</div></div><p style="font-size:11pt;font-family:Calibri,sans-serif;">The final allocable purchase price consists of the following (in thousands):</p><table style="margin-left:auto;border-spacing:0;table-layout:fixed;width:80.0%;border-collapse:separate;margin-right:auto;"><tr style="visibility:collapse;"><td style="width:77.92%;"></td><td style="width:10.1%;"></td><td style="width:1%;"></td><td style="width:9.98%;"></td><td style="width:1.0%;"></td></tr><tr style="height:10pt;background-color:#fffff;word-break:break-word;"><td style="width:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">Class A Common Stock</td><td style="width:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">(1)</td><td style="width:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">27,646,255</td><td style="width:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">851,202</td><td style="width:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">85,120,200</td></tr><tr style="height:10pt;background-color:#fffff;word-break:break-word;"><td style="width:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">Consideration Warrants</td><td style="width:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">(3)</td><td style="width:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">27,646,255</td><td style="width:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">851,202</td><td style="width:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">85,120,200</td></tr><tr style="height:10pt;background-color:#fffff;word-break:break-word;"><td style="width:10pt;margin-top:0;font

align:bottom;border-bottom:0.5pt solid #000000;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"><p></p><td style="white-space:pre-wrap;vertical-align:bottom;border-bottom:0.5pt solid #000000;text-align:center;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:right;">7,434</p></td><td style="white-space:nowrap;vertical-align:bottom;border-bottom:0.5pt solid #ffffff03;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td></tr><tr style="height:10pt;background-color:#fffff;word-break:break-word;"><td style="text-indent:5pt;white-space:pre-wrap;vertical-align:top;"><p style="margin-left:5pt;text-indent:0;font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">Total allocable purchase price</p></td><td style="white-space:pre-wrap;vertical-align:bottom;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td><td style="white-space:nowrap;vertical-align:bottom;border-bottom:2.25pt double #000000;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">\$</p></td><td style="white-space:pre-wrap;vertical-align:bottom;border-bottom:2.25pt double #000000;text-align:center;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:right;">397,251</p></td><td style="white-space:nowrap;vertical-align:bottom;border-bottom:2.25pt double #ffffff03;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td></tr><tr style="background-color:#fffff;word-break:break-word;white-space:pre-wrap;"><td style="vertical-align:bottom;"><p style="font-size:11pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td><td style="vertical-align:bottom;"><p style="font-size:11pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td><td colspan="2" style="vertical-align:bottom;"><p style="font-size:11pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td></tr></table><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;"></p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">(1)27,646,255 shares issued at3.23 per share as of the Acquisition Date.(2)851,202 shares issued at as converted value of323 per share.(3)(3)2,893,731 warrants issued at the estimated fair value of0.2641 per warrant.(4)The portion recorded in consideration transferred represents the fair value of replacement awards (Assumed Options & Assumed RSUs) attributable to precombination services.<div><p style="text-indent:6.667%;font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">We accounted for the transaction as an asset acquisition since the lead asset, taletrectinib, represented substantially all of the fair value of the gross assets acquired, which exclude cash acquired. At the Acquisition Date, we recorded a425.1 million charge representing an acquired in-process research and development asset with no alternative future use in acquired in-process research and development expenses, of which the425.1 million net cash consideration is presented as a cash outflow from operating activities. In connection with this acquisition, we recorded450.9 million of assets acquired primarily consisting of cash and investments. Liabilities assumed were approximately53.7 million.The final allocable purchase price consists of the following (in thousands):</p><table style="margin-left:auto;border-spacing:0;table-layout:fixed;width:80.0%;border-collapse:separate;margin-right:auto;"><tr style="visibility:collapse;"><td style="width:77.92%"><td style="width:1.1%"><td style="width:1.1%"><td style="width:1.1%"><td style="width:1.98%"></td></tr><tr><td style="width:1%"><td style="height:10pt;background-color:#effcf;word-break:break-word;"><td style="white-space:pre-wrap;vertical-align:top;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">Class A Common Stock(1)</p></td><td style="white-space:pre-wrap;vertical-align:bottom;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td><td style="white-space:nowrap;vertical-align:bottom;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">\$</p></td><td style="white-space:pre-wrap;vertical-align:bottom;text-align:center;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:right;">274,938</p></td><td style="white-space:nowrap;vertical-align:bottom;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td></tr><tr><td style="height:10pt;background-color:#effcf;word-break:break-word;"><td style="white-space:pre-wrap;vertical-align:top;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">Series A Non-Voting Convertible Preferred Stock(2)</p></td><td style="white-space:pre-wrap;vertical-align:bottom;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td><td style="white-space:nowrap;vertical-align:bottom;text-align:center;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:right;">89,297</p></td><td

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size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">In October 2023, AnHeart entered into an agreement with NK, granting NK a sub-licensable, royalty-bearing, exclusive right and license to commercialize AB-106 in Japan (the "NK Territory"). AnHeart is responsible for funding ongoing clinical trials of AB-106 in the NK Territory, with NK responsible for funding regulatory submissions in the NK Territory. The Company also granted NK a sub-licensable, royalty-bearing, exclusive right and license to research, develop and commercialize any new indications of AB-106 in the NK Territory ("NK New Indication Right").</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">AnHeart determined that the license represents a single performance obligation. The R&D services represent a material promise and were determined to be a separate performance obligation at the outset of the agreement as the promise is distinct and has standalone value to NK. They concluded that, at the inception of the agreement, the NK New Indication Right did not constitute a material right.</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The portion of the transaction price allocated to the R&D services was deferred and is being recognized as R&D services revenue using an input method. Estimated costs to complete are reassessed on a periodic basis and any updates to the revenue earned are recognized on a prospective basis. The Company recognized R&D service revenue of\$0.3 million for the three months ended June 30, 2024. As of June 30, 2024, the accounts receivable of NK was \$0.1 million millionContract assets and contract liabilities</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">When the Company satisfies its performance obligations by providing services to a customer before the customer pays consideration and before payment is due, the Company recognizes its rights to consideration as a contract asset.</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The Company didn't recognize any contract assets as of June 30, 2024.</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">When a customer pays consideration before the Company provide services, the Company records its obligation as a contract liability.The Company expects to recognize all of this balance as revenue over the next 4 years.</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The contract liabilities of the Company as of June 30, 2024 is listed in the table below.<table style="margin-left:auto;border-spacing:0;table-layout:fixed;width:90.0%;border-collapse:separate;margin-right:auto;"><tr style="visibility:collapse;"><td style="width:80.22%;"></td><td style="width:1.48%;"></td><td style="width:1%;"></td><td style="width:16.3%;"></td><td style="width:1%;"></td></tr><tr style="height:8pt;background-color:#fffff;word-break:break-word;white-space:pre-wrap;"><td style="vertical-align:top;"></td><td style="vertical-align:bottom;"></td><td colspan="2" style="vertical-align:bottom;border-bottom:0.5pt solid #000000;text-align:left;">June 30, 2024</td><td style="vertical-align:bottom;border-bottom:0.5pt solid #fffff0;text-align:left;"></td></tr><tr style="height:8pt;background-color:#fffff;word-break:break-word;white-space:pre-wrap;"><td style="vertical-align:top;"></td><td style="vertical-align:bottom;"></td><td colspan="2" style="vertical-align:bottom;text-align:left;">June 30, 2024</td><td style="vertical-align:bottom;text-align:left;"></td></tr><tr style="height:10pt;background-color:#ff0f;word-break:break-word;"><td style="vertical-align:top;"></td><td style="vertical-align:bottom;"></td><td colspan="2" style="vertical-align:bottom;text-align:left;"></td><td style="vertical-align:bottom;text-align:left;"></td></tr><tr style="height:10pt;background-color:#ff0f;word-break:break-word;"><td style="vertical-align:top;"></td><td style="vertical-align:bottom;"></td><td colspan="2" style="vertical-align:bottom;text-align:left;"></td><td style="vertical-align:bottom;text-align:left;"></td></tr><tr style="height:10pt;background-color:#ff0f;word-break:break-word;"><td style="vertical-align:top;"></td><td style="vertical-align:bottom;"></td><td colspan="2" style="vertical-align:bottom;text-align:left;"></td><td style="vertical-align:bottom;text-align:left;"></td></tr><tr style="height:10pt;background-color:#ff0f;word-break:break-word;"><td style="vertical-align:top;"></td>&

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<content>% per annum.<p style="font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">On April 4, 2023, AnHeart entered into a Loan and Security Agreement ("2023 SSVB Agreement") with Shanghai Pudong Development Bank in Silicon Valley for up to40.0million RMB or equivalent in optional currency USD term loans ("SSVB Loan"). The SSVB Loan consists of a short-term working capital loan of20million RMB and a long-term loan of20million RMB, which matures onApril 4, 2024andApril 4, 2025respectively, at which time all outstanding balances are due. Draws on the line of credit for the short-term loans are payable on the maturity date of the SSVB Loan. Draws on the line of credit for the long-term loans are payable in equal instalments monthly over the shorter of 24 months or from the date the draw took place till the SSVB loan matures. Outstanding balances will bear interest at a fixed rate of4.25%.As of June 30, 2024, the outstanding line of credit balance net of repayments was\$1.8miHion.For the three months ended June 30, 2024, the Company had repaid \$0.7million.70000000700000056000000.0394000000020000000200000002024-04-042025-04-040.04251800000700000Our principal executive office is located in New York, New York, where we lease approximately7,900square feet of office space under a lease that terminates in2027with an option for the Company toextendthe lease for an additionalfive yearswhich is not reasonably assured of exercise. We also occupy approximately25,139square feet of office space in San Francisco, California, under a lease that terminates in2025and a total of approximately1,582square meters of office space in the People's Republic of China, in the cities of Beijing, Guangzhou, Hangzhou, Shanghai and Yantai, under leases that terminate in 2024 through 2026.</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;"></p>Operating lease expense was\$0.5million\$</

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Although the convertible preferred stock are participating securities, such securities do not participate in net losses and therefore do not impact the Company's net loss from continuing operations per share calculation as of June 30, 2024.</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;"><p>Basic loss per share is computed by dividing net loss by the weighted-average number of shares of Class A and Class B Common Stock outstanding. Diluted loss per share reflects the potential dilution that could occur if the stock options to issue Common</p><p><p style="font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;"><p>Stock were exercised. The Company had a net loss in all periods presented thus the dilutive net loss per common share is the same as the basic net loss per common share as the effect of any options or conversions is anti-dilutive.</p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;"><p>The earnings per share amounts are the same for the different classes of Common Stock because the holders of each class are legally entitled to equal per share distributions whether through dividends or liquidation.</p><p><div style="font-size:10pt;font-family:Times New Roman;"><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;"><p>The following securities outstanding at June 30, 2024 and 2023 have been excluded from the calculation of weighted average shares outstanding:</p><p><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></table><table style="margin-left:auto;border-spacing:0;table-layout:fixed;width:80.0%;border-collapse:separate;margin-right:auto;"><tr style="visibility:collapse;"><td style="width:67.313%"></td><td style="width:1.02%"></td><td style="width:1%"></td><td style="width:13.323%"></td><td style="width:1%"></td><td style="width:1.02%"></td><td style="width:1%"></td><td style="width:13.323%"></td><td style="width:1%"></td></tr><tr style="height:8pt;background-color:#fffff0;word-break:break-word;white-space:pre-wrap;text-align:left;"><td style="vertical-align:bottom;"><p style="font-size:8pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:center;"></p></td><td style="vertical-align:bottom;"><p style="font-size:8pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:center;"></p></td><td colspan="6"><p style="vertical-align:bottom;border-bottom:0.5pt solid #000000;"><p style="font-size:8pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:center;">As of June 30,</p></td><td style="vertical-align:bottom;border-bottom:0.5pt solid #fffff03;"><p style="font-size:8pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:center;"></p></td></tr><tr style="height:8pt;background-color:#fffff0;word-break:break-word;white-space:pre-wrap;text-align:left;"><td style="vertical-align:bottom;"><p style="font-size:8pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:center;"></p></td><td style="font-size:8pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:center;"></td><td colspan="6"><p style="vertical-align:bottom;border-bottom:0.5pt solid #000000;"><p style="font-size:8pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:center;">2024</p></td><td style="vertical-align:bottom;border-bottom:0.5pt solid #fffff03;"><p style="font-size:8pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:center;"></p></td></tr><tr style="height:8pt;background-color:#fffff0;word-break:break-word;white-space:pre-wrap;text-align:left;"><td style="vertical-align:bottom;"><p style="font-size:8pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:center;"></p></td><td style="font-size:8pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:center;"></td><td colspan="6"><p style="vertical-align:bottom;border-bottom:0.5pt solid #000000;"><p style="font-size:8pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:center;">2023</p></td><td style="vertical-align:bottom;border-bottom:0.5pt solid #fffff03;"><p style="font-size:8pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:center;"></p></td></tr><tr style="height:10pt;background-color:#ff0ff0;word-break:break-word;"><td style="white-space:pre-wrap;vertical-align:bottom;"><p style="font-size:10pt;margin-top:0;font-family:Times New 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style="height:10pt;background-color:#fffff0;word-break:break-word;"><td style="white-space:pre-wrap;vertical-align:bottom;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td><td style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td><td colspan="6"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td><td style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td></tr><tr style="height:10pt;background-color:#fffff0;word-break:break-word;"><td style="white-space:pre-wrap;vertical-align:bottom;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td><td style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td><td colspan="6"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td><td style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"></p></td></tr><tr style="height:10pt;background-color:#fffff0;word-break:break-word;"&

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Amount reclassified for realized gain on marketable securities

Balance at beginning of period

Currency translation adjustment	
Unrealized gain (loss)	
Total	\$ 195

The following table presents a rollforward of the changes in accumulated other comprehensive income (losses) for the six months ended June 30, 2024 and 2023, which is all attributable to unrealized gains (losses) on available-for-sale securities. All amounts are net of tax.

	2024	2023
Balances at January 1:	\$ 195	\$ 195
Net change during the period:	()	()
Balances at June 30:	\$ 195	\$ 195

space:nowrap;vertical-align:bottom;border-bottom:0.5pt solid #000000;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"><p></td><td style="white-space:pre-wrap;vertical-align:bottom;border-bottom:0.5pt solid #000000;text-align:center;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:right;"><p></td><td style="white-space:nowrap;vertical-align:bottom;border-bottom:0.5pt solid #ffffff03;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"><p></td><tr style="height:10pt;background-color:#ffff0f;word-break:break-word;"><td style="white-space:pre-wrap;vertical-align-top;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;">Balance at end of period<p></td><td style="white-space:pre-wrap;vertical-align:bottom;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"><p></td><td style="white-space:nowrap;vertical-align:bottom;border-bottom:2.25pt double #000000;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"><p></td><td style="white-space:pre-wrap;vertical-align:bottom;border-bottom:2.25pt double #000000;text-align:center;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:right;"><p></td><td style="white-space:nowrap;vertical-align:bottom;border-bottom:2.25pt double #ffffff03;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"><p></td><td style="white-space:pre-wrap;vertical-align:bottom;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"><p></td><td style="white-space:nowrap;vertical-align:bottom;border-bottom:2.25pt double #000000;"><p style="font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:left;"><p></td><tr><table 181000-52526000-1601100093600060001950001480000-1266000-4395000<p style="font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">10. Stock-Based Compensation<p><p style="font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:left;">The 2021 Equity Incentive Plan<p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">In March 2019, the Company adopted the 2019 Equity Incentive Plan or ("2019 Plan"), which provided for the grant of options, stock appreciation rights, restricted stock, and other stock awards. In January 2021, our board of directors adopted the 2021 Equity Incentive Plan (the "2021 Plan"). The 2021 Plan was approved by our stockholders in February 2021 and became effective immediately upon the Closing Date of the Merger. Shares available for future issuance under the 2019 Plan were canceled.<p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Awards.The 2021 Plan provides for the grant of incentive stock options ("ISOs"), within the meaning of Section 422 of the Code to employees, including employees of any parent or subsidiary, and for the grant of nonstatutory stock options ("NSOs"), stock appreciation rights, restricted stock awards, restricted stock unit awards, performance awards and other forms of awards to employees, directors and consultants, including employees and consultants of our affiliates.<p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Authorized Shares.The maximum number of shares of Class A Common Stock reserved for issuance upon adoption of the 2021 Plan was50,684,047shares of Class A Common Stock.The number of shares of Class A Common Stock reserved for issuance under the 2021 Plan will automatically increase on January 1 of each year, starting on January 1, 2022 through January 1, 2031, in an amount equal to (1)4.0% of the total number of shares of Class A Common Stock and Class B Common Stock outstanding or issuable upon conversion or exercise of outstanding instruments on December 31 of the preceding year, or (2) a lesser number of shares of Class A Common Stock determined by our board of directors prior to the date of the increase.The maximum number of shares of Class A Common Stock that may be issued on the exercise of ISOs under the 2021 Plan is three times the number of shares available for issuance upon the 2021 Plan becoming effective or152,052,141shares.<p><p style="font-size:10pt;margin-top:12pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The Employee Stock Purchase Plan<p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">In January 2021, our board of directors adopted the 2021 Employee Stock Purchase Plan (the "ESPP"). The ESPP was approved by our stockholders in February 2021 and became effective immediately upon the Closing Date of the Merger.<p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">Share Reserve. The maximum number of shares of Class A Common Stock reserved for issuance upon the adoption of the 2021 ESPP was4,750,354shares of Class A Common Stock.The number of shares of Class A Common Stock reserved for issuance under the 2021 ESPP will automatically increase on January 1st of each year, beginning on January 1, 2022 and continuing through and including January 1, 2031, by1.0%of the total number of shares of Class A Common Stock and Class B Common Stock outstandingorissuable upon conversion or exercise of outstanding instruments on December 31st of the preceding calendar year or such lesser number of shares of Class A Common Stock as determined by our board of directors.Shares subject to purchase rights granted under the 2021 ESPP that terminate without having been exercised in full will not reduce the number of shares available for issuance under the 2021 ESPP.<div style="font-size:12pt;font-family:Times New Roman;"><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">The stock-based compensation expense included in the Company's condensed statement of operations and comprehensive loss for the three and six months ended June 30, 2024 and 2023 is as follows (in thousands):<p><p style="text-indent:6.667%;font-size:10pt;margin-top:0;font-family:Times New Roman;margin-bottom:0;text-align:justify;"><table style="margin-left:auto;border-spacing:0;table-layout:fixed;width:90.0%;border-collapse:separate;margin-right:auto;">

[illegible]

[illegible]

[illegible]

[illegible]

UNITED STATES SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 20549

Form 10-K

For the fiscal year ended June 30, 2024

1. Business

Our company is a blank check company. We have not yet identified any business to acquire. We are currently in the process of identifying potential business opportunities and evaluating them. We have not yet entered into any definitive agreements with any potential business targets. We are currently in the process of identifying potential business opportunities and evaluating them. We have not yet entered into any definitive agreements with any potential business targets.

2. Description of Business

Our company is a blank check company. We have not yet identified any business to acquire. We are currently in the process of identifying potential business opportunities and evaluating them. We have not yet entered into any definitive agreements with any potential business targets. We are currently in the process of identifying potential business opportunities and evaluating them. We have not yet entered into any definitive agreements with any potential business targets.

3. Legal Proceedings

There are no legal proceedings pending against our company or involving our company.

4. Risk Factors

There are no risk factors associated with our company.

5. Management and Directors

Our management and directors are as follows:

Name	Position
John Doe	Chief Executive Officer
Jane Smith	Chief Financial Officer
Bob Johnson	Chief Operating Officer
Alice Brown	Chief Marketing Officer
Charlie White	Chief Technology Officer

6. Compensation

Our compensation policy is as follows:

Position	Salary	Bonus
Chief Executive Officer	\$1,000,000	\$500,000
Chief Financial Officer	\$500,000	\$250,000
Chief Operating Officer	\$400,000	\$200,000
Chief Marketing Officer	\$300,000	\$150,000
Chief Technology Officer	\$300,000	\$150,000

7. Financial Information

Our financial information is as follows:

Item	Amount
Revenue	\$0
Expenses	\$0
Net Income	\$0
Assets	\$0
Liabilities	\$0
Equity	\$0

8. Financial Statements

Our financial statements are as follows:

Item	Amount
Revenue	\$0
Expenses	\$0
Net Income	\$0
Assets	\$0
Liabilities	\$0
Equity	\$0

9. Additional Information

Our additional information is as follows:

Item	Amount
Revenue	\$0
Expenses	\$0
Net Income	\$0
Assets	\$0
Liabilities	\$0
Equity	\$0

10. Signatures

Our signatures are as follows:

Name	Signature
John Doe	[Signature]
Jane Smith	[Signature]
Bob Johnson	[Signature]
Alice Brown	[Signature]
Charlie White	[Signature]

[illegible]

[illegible]

[illegible]

[illegible]

content;">June 21, 2024, >David Hanley, the Company's >Chief Technical Operations Officer,> entered into a trading plan intended to satisfy the affirmative defense conditions of >(c) under the Exchange Act, providing for the sale of up to 500,000> shares of the Company's Common Stock. Pursuant to this plan, Mr. Hanley may sell shares beginning October 21, 2024 and ending November 1, 2025.> </p><p style="text-indent:6.667%;font-size:10pt;margin-top:6pt;font-family:Times New Roman;margin-bottom:0;text-align:justify;">On >June 21, 2024,>>Gary Hattersley,>the Company's >Chief Scientific Officer,> entered into a trading plan intended to satisfy the affirmative defense conditions of >Rule 10b5-1>(c) under the Exchange Act, providing for the sale of up to 398,935> shares of the Company's Common Stock. Pursuant to this plan, Mr. Hanley may sell shares beginning October 21, 2024 and ending November 1, 2026.> </p><p>>June 21, 2024>Kerry Wentworth>Chief Regulatory Officer>true303920>June 21, 2024>David Hanley>Chief Technical Operations Officer>true500000>June 21, 2024>Gary Hattersley>Chief Scientific Officer>true398935>27,646,255>shares issued at \$3.23 per share as of the Acquisition Date.>851,202>shares issued at as converted value of \$323 per share.>2,893,731>warrants issued at the estimated fair value of \$0.2641 per warrant. The portion recorded in consideration transferred represents the fair value of replacement awards (Assumed Options & RSTs) attributable to precombination services.</p></div>