â€‹â€‹UNITED STATESSECURITIES AND EXCHANGE COMMISSIONWashington, D.C. 20549FORMÂ 6-KREPORT OF FOREIGN ISSUERPURSUANT TO RULEÂ 13a-16 OR 15d-16OF THE SECURITIES EXCHANGE ACT OF 1934For theÂ month of December 2024(Commission File No.Â 001-40241)LAVA Therapeutics N.V.(Translation of registrantâ€™s name into English)Yalelaan 623584 CM Utrecht, The Netherlands(Address of principal executive office)Indicate by check mark whether the registrant files or will file annual reports under cover of FormÂ 20-F or FormÂ 40-F.FormÂ 20-FÂ Â â˜’ FormÂ 40-FÂ Â â˜â€‹â€‹â€‹â€‹â€‹LAVA Therapeutics, N.V.Exhibits 99.1 and 99.2 to this Report on FormÂ 6-K shall be deemed to be incorporated by reference into the registration statement on FormÂ S-8 (File no. 333-256655) and registration statement on Form F-3 (File no. 333-264246) of LAVA Therapeutics N.V. (including any prospectuses forming a part of such registration statements) and to be a part thereof from the date on which this report is filed, to the extent not superseded by documents or reports subsequently filed or furnished.ExhibitÂ 99.3 to this Report on Form 6-K is being furnished and shall not be deemed â€œfiledâ€ for purposes of SectionÂ 18 of the Securities Exchange Act of 1934, as amended (Exchange Act), or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act.â€‹RISK FACTORSOur business faces significant risks. You should carefully consider all of the information set forth in this report and in our other filings with the United States Securities and Exchange Commission (SEC). Our business, financial condition, results of operations and growth prospects could be materially adversely affected by any of these risks. This report also contains forward-looking statements that involve risks and uncertainties. Our results could materially differ from those anticipated in these forward-looking statements, as a result of certain factors including the risks described in our other SEC filings.â€‹â€‹Exhibit ListExhibit ListExhibitÂ Â Â Â Description99.1â€‹Unaudited Condensed Consolidated Interim Financial Statements as of and for the Three and Nine Months Ended September 30, 2024 and 202399.2â€‹Managementâ€™s Discussion and Analysis of Financial Condition and Results of Operations as of and for the Three and Nine Months Ended September 30, 2024 and 202399.3â€‹Press Release dated December 10, 2024â€‹â€‹â€‹â€‹â€‹SIGNATURESPursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.â€‹LAVA Therapeutics, N.V.â€‹(Registrant)â€‹â€‹â€‹Date: December 10, 2024By:/s/ Fred Powellâ€‹Fred Powellâ€‹Chief Financial Officerâ€‹Table of ContentsExhibitÂ 99.1LAVA THERAPEUTICS N.V.INDEX TO UNAUDITED CONDENSED CONSOLIDATED INTERIM FINANCIAL STATEMENTSâ€‹PageCondensed Consolidated Interim Statements of Loss and Comprehensive Loss (unaudited)2Condensed Consolidated Interim Statements of Financial Position (unaudited)3Condensed Consolidated Interim Statements of Changes in Equity (unaudited)4Condensed Consolidated Interim Statements of Cash Flows (unaudited)6NotesÂ to the Condensed Consolidated Interim Financial Statements (unaudited)7â€‹â€‹â€‹â€‹Table of ContentsLAVA Therapeutics N.V.Condensed Consolidated Interim Statements of Lossand Comprehensive Loss(in thousands, except share and per share amounts) (unaudited)â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Â Â Â Â Â Â Â Â Â Â Â Three Months Ended Â Â Â Â Nine Months Ended â€‹â€‹â€‹â€‹â€‹SeptemberÂ 30,Â â€‹SeptemberÂ 30,Â â€‹â€‹Â Â Â Â NotesÂ Â Â Â 2024Â Â Â Â 2023Â Â Â Â 2024Â Â Â Â 2023â€‹Revenue:Â Â Â Â â€‹Â Â Â â€‹Â Â Â â€‹Â Â Â â€‹Â Â â€‹Revenue from contracts with customersÂ 5â€‹$ â€”â€‹$ 53â€‹$ 6,992â€‹$ 6,416â€‹Cost of sales of goodsâ€‹5â€‹â€‹ â€”â€‹â€‹ â€”â€‹â€‹ â€”â€‹â€‹ (2,546)â€‹Cost of providing servicesâ€‹5â€‹â€‹ â€”â€‹â€‹ (10)â€‹â€‹ â€”â€‹â€‹ (782)â€‹Gross profitÂ Â Â â€‹Â â€”â€‹Â 43â€‹Â 6,992â€‹Â 3,088â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Operating expenses:Â Â Â â€‹Â Â Â â€‹Â Â Â â€‹Â Â Â â€‹Â Â Â â€‹Research and developmentÂ 6â€‹Â (8,500)â€‹Â (7,912)â€‹Â (20,844)â€‹Â (30,454)â€‹General and administrativeÂ 7â€‹Â (2,785)â€‹Â (2,858)â€‹Â (8,745)â€‹Â (10,445)â€‹Total operating expensesÂ Â Â â€‹Â (11,285)â€‹Â (10,770)â€‹Â (29,589)â€‹Â (40,899)â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Operating lossÂ Â Â â€‹Â (11,285)â€‹Â (10,727)â€‹Â (22,597)â€‹Â (37,811)â€‹Interest income, netÂ â€‹â€‹Â 805â€‹Â 809â€‹Â 2,426â€‹Â 2,124â€‹Foreign currency exchange (loss) gain, netÂ â€‹â€‹Â (1,720)â€‹Â 1,132â€‹Â (723)â€‹Â 429â€‹Total non-operating (loss) incomeÂ Â Â â€‹Â (915)â€‹Â 1,941â€‹Â 1,703â€‹Â 2,553â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Loss before income taxÂ Â Â â€‹Â (12,200)â€‹Â (8,786)â€‹Â (20,894)â€‹Â (35,258)â€‹Income tax expenseÂ â€‹â€‹Â (96)â€‹Â (50)â€‹Â (250)â€‹Â (218)â€‹Loss for the periodÂ Â Â â€‹$ (12,296)â€‹$ (8,836)â€‹$ (21,144)â€‹$ (35,476)â€‹Items that may be reclassified to profit or lossâ€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Foreign currency translation adjustmentÂ â€‹â€‹Â 1,688â€‹Â (1,385)â€‹Â 296â€‹Â (82)â€‹Total comprehensive lossÂ Â Â â€‹$ (10,608)â€‹$ (10,221)â€‹$ (20,848)â€‹$ (35,558)â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Â Â â€‹â€‹â€‹â€‹â€‹â€‹â€‹Loss per share:Â Â Â â€‹Â Â Â â€‹Â Â Â â€‹Â Â Â â€‹Â Â Â â€‹Loss per share, basic and dilutedâ€‹â€‹$ (0.46)â€‹$ (0.34)â€‹$ (0.79)â€‹$ (1.35)â€‹Weighted-average common shares outstanding, basic and dilutedâ€‹â€‹â€‹Â 26,846,006â€‹Â 26,289,087â€‹Â 26,814,113â€‹Â 26,289,087â€‹â€‹â€‹The accompanying notes are an integral part of the unaudited condensed consolidated interim financial statements.â€‹2Table of ContentsLAVA Therapeutics N.V.Condensed Consolidated Interim Statements of Financial Position(in thousands) (unaudited)â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹SeptemberÂ 30,Â â€‹DecemberÂ 31,Â â€‹â€‹Â Â Â Â Â Â Â Â 2024Â Â Â Â 2023AssetsÂ Â â€‹Â Â Â â€‹Â Â Non-current assets:Â Â â€‹Â Â Â â€‹Â Â Property and equipment, netÂ â€‹$ 1,188â€‹$ 1,602Right-of-use assetsÂ â€‹Â 517â€‹Â 892Other non-current assets and security depositsÂ â€‹Â 114â€‹Â 319Total non-current assetsÂ â€‹Â 1,819â€‹Â 2,813Current assets:Â â€‹Â Â Â â€‹Â Â Receivables and otherÂ â€‹Â 536â€‹Â 1,459Prepaid expenses and other current assetsÂ â€‹Â 1,629â€‹Â 1,627VAT receivableÂ â€‹Â 489â€‹Â 240Investmentsâ€‹â€‹â€‹ 51,921â€‹â€‹ 51,340Cash and cash equivalentsÂ â€‹Â 26,963â€‹Â 44,231Total current assetsÂ â€‹Â 81,538â€‹Â 98,897Total assetsÂ â€‹$ 83,357â€‹$ 101,710Equity and LiabilitiesÂ â€‹Â Â Â â€‹Â Â Â Equity:Â â€‹Â Â Â â€‹Â Â Â Share capitalÂ â€‹$ 3,716â€‹$ 3,715Equity-settled employee benefits reserveÂ â€‹Â 14,360â€‹Â 12,005Foreign currency translation reserveÂ â€‹Â (10,603)â€‹Â (10,899)Additional paid-in capitalÂ â€‹Â 194,450â€‹Â 194,424Accumulated deficitÂ â€‹Â (168,930)â€‹Â (148,067)Total equityÂ â€‹Â 32,993â€‹Â 51,178Non-current liabilities:Â â€‹Â Â Â â€‹Â Â Â Deferred revenueÂ â€‹Â 35,000â€‹Â 35,000Lease liabilitiesÂ â€‹Â 164â€‹Â 591Total non-current liabilitiesÂ â€‹Â 35,164â€‹Â 35,591Current liabilities:Â â€‹Â Â Â â€‹Â Â Â Trade payables and otherÂ â€‹Â 2,815â€‹Â 4,446Borrowingsâ€‹â€‹â€‹ 5,756â€‹â€‹ 5,282Lease liabilitiesÂ â€‹Â 370â€‹Â 440Accrued expenses and other current liabilitiesÂ â€‹Â 6,259â€‹Â 4,773Total current liabilitiesÂ â€‹Â 15,200â€‹Â 14,941Total liabilitiesÂ â€‹Â 50,364â€‹Â 50,532Total equity and liabilitiesÂ â€‹$ 83,357â€‹$ 101,710â€‹â€‹â€‹â€‹The accompanying notes are an integral part of the unaudited condensed consolidated interim financial statements.â€‹â€‹â€‹3Table of ContentsLAVA Therapeutics N.V.Condensed Consolidated Interim Statements of Changes in Equity(in thousands, except share amounts) (unaudited)â€‹
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Using its Gammabody platform, the Company is developing a portfolio of novel bispecific antibodies designed to engage and leverage the potency and precision of gamma-delta (gd) T cells to elicit a robust, anti-tumor immune response and improve outcomes for cancer patients. LAVA Therapeutics N.V. was incorporated in 2016 and is headquartered in Utrecht, the Netherlands. In August 2024, we established a wholly owned Australian subsidiary, LAVA Therapeutics (Australia) Pty Ltd. Unless the context otherwise requires, references to the â€œCompany,â€ â€œwe,â€ â€œusâ€ and â€œourâ€ refer to LAVA Therapeutics N.V. and its subsidiaries.In connection with becoming a public company, on MarchÂ 29, 2021, the Company converted from â€œLAVA Therapeutics, B.V.â€ to â€œLAVA Therapeutics N.V.â€ The address of the Companyâ€™s registered office is Yalelaan 62, 3584 CM Utrecht, the Netherlands. The Companyâ€™s common shares are listed for trading under the symbol â€œLVTXâ€ on The Nasdaq Global Select Market.The Company will transition from foreign private issuer to U.S. domestic filer status on January 1, 2025 and will file its annual report for the year ended December 31, 2024 on a Form 10-K. The Company will prepare the financial statements in accordance with U.S. Generally Accepted Accounting Principles (US GAAP) for all comparative periods. â€‹The Audit Committee of the Companyâ€™s Board of Directors approved these unaudited condensed consolidated interim financial statements on December 10, 2024.NoteÂ 2â€”Summary of Significant Accounting PoliciesBasis of preparationThe unaudited condensed consolidated interim financial statements of the Company are prepared in accordance with International Accounting Standard 34, â€œInterim Financial Reportingâ€ as issued by the International Accounting Standards Board. Certain information and disclosures normally included in the consolidated financial statements prepared in accordance with International Financial Reporting Standards (IFRS) have been condensed or omitted. Accordingly, these unaudited condensed consolidated interim financial statements should be read in conjunction with the Companyâ€™s annual consolidated financial statements for theÂ year ended DecemberÂ 31, 2023 and accompanying notes, which have been prepared in accordance with IFRS as issued by the International Accounting Standards Board and are included on Form 20-F filed by the Company on March 20, 2024.The accounting policies applied are consistent with those of the previous financialÂ year. A description of our accounting policies is provided in the Accounting Policies section of the audited consolidated financial statements as of and for theÂ years ended December 31, 2023 and 2022, included on Form 20-F filed by the Company on March 20, 2024.The preparation of financial statements in conformity with IFRS requires the use of certain critical accounting estimates and requires management to exercise its judgment in the process of applying the Companyâ€™s accounting policies. The areas involving a higher degree of judgment or complexity or areas where assumptions and estimates are significant to the unaudited condensed consolidated interim financial statements are disclosed in NoteÂ 3. The interim financial data as of September 30, 2024 and 2023, and for the three and nine months ended September 30, 2024 and 2023, are unaudited. In the opinion of 7Table of Contentsmanagement, the interim financial data includes all adjustments, consisting only of normal recurring adjustments, necessary to a fair statement of the results for the interim periods.License RevenueWe may enter into collaboration and licensing arrangements for research and development, manufacturing, and commercialization activities with counterparties for the development and commercialization of our product candidates. These arrangements may contain multiple components, such as (i) licenses, (ii) research and development activities, and (iii) the manufacturing of certain materials. Payments pursuant to these arrangements may include non-refundable and refundable payments, payments upon the achievement of significant regulatory, development and commercial milestones, sales of products at certain agreed-upon amounts, and royalties on product sales.In determining the appropriate amount of revenue to be recognized as we fulfill our obligations under a collaboration agreement, we perform the following steps: (i) identification of the promised goods or services in the contract; (ii) determination of whether the promised goods or services are performance obligations, including whether they are capable of being distinct; (iii) measurement of the transaction price, including the constraint on variable consideration; (iv) allocation of the transaction price to the performance obligations; and (v) recognition of revenue as we satisfy each performance obligation.We must develop estimates and assumptions that require judgment to determine the underlying stand-alone selling price for each performance obligation, which determines how the transaction price is allocated among the performance obligations. The estimation of the stand-alone selling price may include such estimates as forecasted revenues and costs, development timelines, discount rates and probabilities of regulatory and commercial success. We also apply significant judgment when evaluating whether contractual obligations represent distinct performance obligations, allocating transaction price to performance obligations within a contract, determining when performance obligations have been met, assessing the recognition and future reversal of variable consideration and determining and applying appropriate methods of measuring progress for performance obligations satisfied

over time.Revision of Immaterial Misstatements: Statement of Cash Flow Presentationâ€‹ During the third quarter of 2024, the Company identified a misstatement in the presentation of condensed consolidated interim statements of cash flows. Â The misstatement relates to the cash flow presentation of accrued interest related to the Rijksdienst voor Ondernemend Nederland (RVO) debt agreement. Under the terms of the RVO agreement, interest accrues over the life of the loan and is payable at the end of the loan term. Under IFRS, accrued interest on debt arrangements is a non-cash adjustment to net loss in cash flows from operating activities. The Company incorrectly classified the activity as a cash inflow from borrowings in cash flows from financing activities in its historical financial statements.Â As a result of this misstatement the Company revised the accrued interest on borrowings (previously called â€œProceeds from borrowingsâ€) of $0.3 million for the period ended September 30, 2023, from financing activities to non-cash adjustments in operating activities. The Company made similar adjustments in the statements of cash flows for the year ended December 31, 2023, three month ended March 31, 2023, six months ended June 30, 2023, three months ended March 31, 2024 and six months ended June 30, 2024 of $0.5 million, $0.1 million, $0.2 million, $0.1 million, and $0.3 million, respectively. There was no impact on the Consolidated Statements of Loss and Comprehensive Loss, Consolidated Statements of Financial Position, or Consolidated Statements of Changes in Equity for any period presented.The Company has concluded that the revision in the historical condensed consolidated interim statements of cash flows is not material, both individually and in aggregate, for any prior period.Â â€‹8Table of ContentsGoing ConcernThrough September 30, 2024, the Company has funded its operations with proceeds from sales of equity, collaboration and licensing agreements, government grants and borrowings under various agreements. Since its inception, the Company has incurred net losses. The Dutch Research and Development Act (WBSO) provides compensation for a part of research and development wages and other costs through a reduction in payroll taxes. WBSO grant amounts are offset against wages and salaries and included in research and development expenses in the condensed consolidated interim statements of loss and comprehensive loss.As of SeptemberÂ 30, 2024, the Company had an accumulated deficit of $168.9Â million. The Company expects to continue to generate operating losses in the foreseeable future. The Company expects that its cash, cash equivalents and investments of $78.9 million as of September 30, 2024, will be sufficient to fund its operating expenses and capital expenditure requirements for at least 12Â months following the issuance of these condensed consolidated interim financial statements. Accordingly, the condensed consolidated interim financial statements have been prepared on a going-concern basis.Until we can generate sufficient product revenue to finance our cash requirements, which we may never do, we expect to finance our future cash needs through a combination of public or private equity offerings, debt financings, collaborations, strategic alliances, licensing arrangements and other marketing or distribution arrangements. Disruptions in the financial markets, in general, may make equity and debt financing more difficult to obtain and may have a material adverse effect on our ability to meet our fundraising needs. If we are unable to obtain sufficient funding in a timely manner or on commercially acceptable terms, we may have to delay, reduce the scope of, or eliminate one or more of our operating activities and consider other cost-reduction initiatives, such as downsizing our operations or withholding initiation or expansion of clinical trials or research. In addition, in the event we are not able to generate sufficient funds, we may be unable to continue as a going concern, and our business, financial condition and/or results of operations could be materially and adversely affected and could reduce the price of our common shares and we may ultimately go into insolvency. In addition, any perceived or actual inability by us to finance our clinical development activities and other business activities may cause the market price of our common shares to decline.Cash and cash equivalentsCash and cash equivalents in the condensed consolidated interim statements of financial position are comprised of cash at banks and short-term deposits with a maturity of threeÂ months or less, which are subject to an insignificant risk of changes in value. Our cash and cash equivalents are held in multiple currencies, primarily in the Euro and United States (U.S.) dollar. Accordingly, our cash balances may be exposed to foreign currency exchange risk.For the purposes of the condensed consolidated interim statements of cash flows, cash and cash equivalents consist of cash and short-term deposits, as defined above, net of outstanding bank overdrafts.InvestmentsAs of SeptemberÂ 30, 2024, we have determined that we have the intent and ability to hold all investments in debt securities until maturity. Accordingly, all investments are recorded at amortized cost on our condensed consolidated interim statements of financial position, with the amortization of bond premiums or discounts and earned interest income recorded in our condensed consolidated interim statements of loss and other comprehensive loss.9Table of ContentsFinancial instruments(i)Financial assetsThe Companyâ€™s financial assets are comprised of cash and cash equivalents, investments, trade and other receivables, security deposits, other current and non-current assets. All financial assets are recognized initially at fair value plus transaction costs that are attributable to the acquisition of the financial asset, and are held for standard working capital purposes. These financial assets are subsequently measured at amortized cost, which in general, approximates to the fair value. Purchases and sales of financial assets are recognized on the settlement date or the date that the Company receives or delivers the asset. The Company classifies its financial assets primarily as cash and cash equivalents and receivables. Receivables are non-derivative financial assets, with fixed or determinable payments that are not quoted in an active market. They are included in current assets.Financial assets are derecognized when the rights to receive cash flows from the asset have expired, or the Company has transferred its rights to receive cash flows from the asset or has assumed an obligation to pay the received cash flows in full.(ii)Financial liabilitiesThe Companyâ€™s financial liabilities are comprised of trade and other payables, lease liabilities, and borrowings. All financial liabilities are recognized initially at fair value, adjusted for transaction costs.After initial recognition, borrowings are subsequently measured at amortized cost using the effective interest method, minus transaction costs that are directly attributable to the financial liability. The effective interest method amortization is included in finance costs in the condensed consolidated interim statements of loss and other comprehensive loss.Payables and borrowings are classified as current liabilities unless the Company has an unconditional right to defer settlement of the liability for at least 12Â months after the reporting date.Financial liabilities are derecognized when the obligation under the liability is discharged, canceled, or expires.(iii)Fair value measurementsThe Company does not hold any financial assets and financial liabilities other than those measured at amortized cost, as its business model is such that the Company has the intent to hold these instruments for the sole purpose of collecting contractual cash flows, and the contractual terms give rise to cash flows that are solely for payments of principal and interest. Management assessed that the carrying values of the Companyâ€™s financial assets and financial liabilities measured at amortized cost are a reasonable approximation of their fair values.NoteÂ 3â€”Material Accounting Judgments, Estimates and AssumptionsIn the application of our accounting policies, the Company is required to make judgments, estimates and assumptions about the carrying amounts of assets and liabilities that are not readily apparent from other sources. The estimates and associated assumptions are based on historical experience and other factors that are considered to be relevant. Actual results may differ from these estimates.The estimates and underlying assumptions are reviewed on an ongoing basis. Revisions to accounting estimates are recognized in the period in which the estimate is revised if the revision affects only that period, or in the period of the revision and future periods if the revision affects both current and future periods.10Table of ContentsCritical judgments made in the process of applying our accounting policies which have the most significant effect on the amounts recognized in our unaudited condensed consolidated interim financial statements relate to revenue recognition, share-based payments, accruals for clinical trial expenses, lease accounting, and to our research and license agreements.The key sources of estimation uncertainty that have a significant risk of causing a material adjustment to the carrying amount of assets and liabilities within the next financialÂ year, primarily relate to recognition of accruals for manufacturing and clinical trial activities. No material adjustments to accruals have been recognized during the first nineÂ months of 2024 or 2023, due to conditions that existed as of DecemberÂ 31, 2023 or 2022, respectively. Additionally, there have been no changes to the application of significant accounting estimates, and no impairment losses have been recognized during the first nineÂ months of 2024 or 2023.New standards, interpretations and amendmentsThere were no new standards, interpretations, or amendments that became effective in the current reporting period which had an impact on the unaudited condensed consolidated interim financial statements.â€‹The unaudited condensed consolidated interim financial statements do not include all disclosures for material accounting judgments, estimates and assumptions that are required in the annual consolidated financial statements and should be read in conjunction with the Companyâ€™s audited consolidated financial statements as of, and for theÂ years ended, DecemberÂ 31, 2023 and 2022.NoteÂ 4â€”Equity The share capital of LAVA Therapeutics N.V. consisted of 26,298,665 issued and outstanding common shares at a nominal value of $0.14 per share as of September 30, 2024.â€‹NoteÂ 5â€”Revenue and cost of salesâ€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Three Months Ended SeptemberÂ 30,Â â€‹Nine Months Ended SeptemberÂ 30,Â (in thousands)Â Â Â Â 2024Â Â Â Â 2023Â Â Â Â 2024Â Â Â Â 2023Pfizer Inc. - Pfizer Agreement - Milestonesâ€‹$ â€”â€‹$ â€”â€‹$ 6,960â€‹$ â€”Pfizer Inc. - Pfizer Agreement - Other activitiesâ€‹â€‹ â€”â€‹â€‹ 27â€‹â€‹ 24â€‹â€‹ 3,422Pfizer Inc. - Additional servicesâ€‹â€‹ â€”â€‹â€‹ 26â€‹â€‹ 8â€‹â€‹ 504Johnson & Johnson Agreement - Milestonesâ€‹Â â€”â€‹Â â€”â€‹Â â€”â€‹Â 2,490Total revenue from contracts with customersâ€‹$ â€”â€‹$ 53â€‹$ 6,992â€‹$ 6,416â€‹Pfizer AgreementIn September 2022, the Company entered the Pfizer Agreement to develop, manufacture and commercialize EGFRd2 (PF-08046052/formerly LAVA-1223), an advanced preclinical asset that utilizes LAVAâ€™s proprietary Gammabody technology to target EGFR-expressing solid tumors. Under the terms of the Pfizer agreement, it received a $50.0 million nonrefundable upfront payment in October 2022 and could receive up to approximately $650.0 million in potential development, regulatory and commercial milestones, and royalties ranging from high single-digit to mid-teen percentages on future sales. The Pfizer agreement also provided Pfizer with the opportunity to exclusively negotiate rights to apply our proprietary Gammabody 11Table of Contentsplatform on up to two additional tumor targets, which rights have expired. In March 2024, Pfizer achieved a clinical development milestone for EGFRd2 (PF-08046052), resulting in the first milestone payment of $7.0 million to the Company under the Pfizer Agreement. This payment was recognized as milestone revenue during the nine months ended September 30, 2024.The Company is entitled to receive tiered royalties based on commercial sales levels from high single-digit to mid-teen percentages of net sales of licensed products. Pfizer has also granted it a one-time option to obtain increased royalties if it exercises a buy-up option within a certain amount of time from certain key early clinical data becoming available for the first licensed product. The Company has a specified period of time after notice of such buy-up option to pay Pfizer a one-time fee of $35.0 million (buy-up fee). In the event the Company exercises the buy-up option and pays the buy-up fee, it is entitled to receive increased future royalty percentages to a range of low double-digit to high teen percentages on future sales, and certain future milestones will be decreased by 30%. The deferred revenue balance related to the buy-up option is considered as a monetary item. Royalties are payable on a licensed product-by-licensed product and country-by-country basis beginning with the first commercial sale of such licensed product in such country of sale and expiring ten years after such sale, subject to specified and capped reductions for the market entry of biosimilar products, loss of patent coverage of licensed products, and for payments owed to third parties for additional rights necessary to commercialize licensed products in the territory.Under the Pfizer Agreement, the Company is also entitled to receive reimbursement of up to $6.5 million for certain agreed to research, manufacturing and supply activities, as well as the transfer of all manufacturing-related know-how and materials, including all CMC documentation, data and processes, to enable the manufacture of licensed compounds and products by Pfizer. During the three months ended September 30, 2024, the Company did not recognize revenue under the Pfizer Agreement.The Company determined that the Pfizer Agreement and the research, manufacturing and supply activities and materials transfer fall within the scope of IFRS 15, Revenue from Contracts with Customers (IFRS 15). In calculating the transaction price, it determined the following four performance obligations under the agreement: (i) provide exclusive license; (ii) provide manufacturing technology transfer activities; (iii) provide initial drug supply; and (iv) research activities, including data and support for regulatory submission.The Company allocated the transaction price to the performance obligations as of September 30, 2024 and 2023 as follows:â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Revenue Recognizedâ€‹Revenue Recognizedâ€‹Cumulative Revenueâ€‹â€‹Transaction â€‹for the quarter endedâ€‹for the nine months endedâ€‹Recognized(in thousands)Â Â Â Â PriceÂ Â Â Â September 30, 2024Â Â Â Â September 30, 2024Â Â Â Â as of September 30, 2024License â€‹$ 50,000â€‹$ â€”â€‹$ â€”â€‹$ 15,165Manufacturing technology transfer activitiesâ€‹â€‹ 2,167â€‹â€‹ â€”â€‹â€‹ 24â€‹â€‹ 2,296Initial supplyâ€‹â€‹ 3,583â€‹â€‹ â€”â€‹â€‹ â€”â€‹â€‹ 3,443Research activitiesâ€‹â€‹ 750â€‹â€‹ â€”â€‹â€‹ â€”â€‹â€‹ 687Buy-up fee (*)â€‹â€‹ (35,000)â€‹â€‹ â€”â€‹â€‹ â€”â€‹â€‹ â€”â€‹â€‹$ 21,500â€‹$ â€”â€‹$ 24â€‹$ 21,591â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹(*) Buy-up fee remains deferred until option expires or is exercisedâ€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹12Table of Contentsâ€‹â€‹â€‹â€‹â€‹Revenue Recognizedâ€‹Revenue Recognizedâ€‹Cumulative Revenueâ€‹â€‹Transactionâ€‹for the quarter endedâ€‹for the nine months endedâ€‹Recognized(in thousands)Â Â Â Â Priceâ€‹September 30, 2023Â Â Â Â September 30, 2023Â Â Â Â as of September 30, 2023Licenseâ€‹$ 50,000â€‹$ â€”â€‹$ â€”â€‹$ 15,165Manufacturing technology transfer activities â€‹â€‹ 2,167â€‹â€‹ 12â€‹â€‹ 112â€‹â€‹ 2,185Initial supplyâ€‹â€‹ 3,583â€‹â€‹ 13â€‹â€‹ 3,287â€‹â€‹ 3,287Research activitiesâ€‹â€‹ 750â€‹â€‹ 2â€‹â€‹ 23â€‹â€‹ 686Buy-up fee (*)â€‹â€‹ (35,000)â€‹â€‹ â€”â€‹â€‹ â€”â€‹â€‹ â€”â€‹â€‹$ 21,500â€‹$ 27â€‹$ 3,422â€‹$ 21,323â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹(*) Buy-up fee remains deferred until option expires or is exercisedâ€‹â€‹â€‹â€‹â€‹â€‹â€‹For the nine months ended September 30, 2023, revenue primarily relates to the initial drug supply provided to Pfizer. Of the initial supply revenue of $3.3 million recognized during the nine months ended September 30, 2023, $0.8 million related to services provided in relation to stability studies, and $2.5 million related to the cost of drug supply provided to Pfizer. Johnson & Johnson (J&J) AgreementMilestone paymentNo milestone revenue was recognized in the three and nine months ended September 30, 2024 under the J&J Agreement (formerly the â€œJanssen Biotech, Inc.â€ agreement). In May 2023, a milestone payment of $2.5 million from J&J was triggered under the terms of the J&J AgreementÂ following the selection of a candidate novel bispecific antibody to engage gamma-delta T cells to an undisclosed tumor associated antigen for the treatment of cancer. Efforts are underway to advance the candidate towards the clinic. This milestone payment was recognized as revenue in the three months ended June 30, 2023 and the payment of the $2.5 million milestone was received in July 2023. NoteÂ 6â€”Research and Development ExpensesResearch and development expenses were as follows:â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Three Months Ended SeptemberÂ 30,Â â€‹Nine Months Ended SeptemberÂ 30,Â â€‹(in thousands)Â Â Â Â 2024Â Â Â Â 2023Â Â Â Â 2024Â Â Â Â 2023Â Â Pre-clinical and clinical trial expensesâ€‹$ 5,722â€‹$ 4,899â€‹$ 12,836â€‹$ 20,542â€‹Personnel-related expensesâ€‹â€‹ 1,304â€‹â€‹ 1,763â€‹â€‹ 3,830â€‹â€‹ 5,346â€‹Research and development activities expensesâ€‹Â 678â€‹Â 325â€‹Â 2,024â€‹Â 1,297â€‹Facilities and other research and development expensesâ€‹Â 551â€‹Â 622â€‹Â 1,192â€‹Â 1,842â€‹Share-based compensation expenseâ€‹â€‹ 245â€‹â€‹ 303â€‹â€‹ 962â€‹â€‹ 1,427â€‹â€‹â€‹$ 8,500â€‹$ 7,912â€‹$ 20,844â€‹$ 30,454â€‹â€‹â€‹â€‹13Table of ContentsNoteÂ 7â€”General and Administrative ExpensesGeneral and administrative expenses were as follows:â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Three Months Ended SeptemberÂ 30,Â â€‹Nine Months Ended SeptemberÂ 30,Â (in thousands)Â Â Â Â 2024Â Â Â Â 2023Â Â Â Â 2024Â Â Â Â 2023Personnel-related expensesâ€‹$ 976â€‹$ 742â€‹$ 2,937â€‹$ 2,932Professional and consultant feesâ€‹Â 792â€‹Â 887â€‹Â 2,356â€‹Â 2,667Insurance, facilities, fees and other related costsâ€‹Â 542â€‹Â 608â€‹Â 1,778â€‹Â 2,159Share-based compensation expenseâ€‹â€‹ 475â€‹â€‹ 621â€‹â€‹ 1,674â€‹â€‹ 2,687â€‹â€‹$ 2,785â€‹$ 2,858â€‹$ 8,745â€‹$ 10,445â€‹â€‹NoteÂ 8â€”Share-based AwardsAs of MarchÂ 25, 2021, the 2018 Stock Option Plan and the 2020 U.S. Stock Option Plan ceased to have any future shares available, and the Company established the 2021 Long-Term Incentive Option Plan (the Plan) for all its employees, members of the Board of Directors and select external consultants.Stock OptionsThere were 6,584,726 stock options outstanding as of September 30, 2024, at a weighted-average exercise price of $3.31 per share. Of these outstanding options 3,592,462 options were vested and exercisable at a weighted-average exercise price of $3.63 per share. During the threeÂ months ended September 30, 2024, 3,150 options were granted to employees at a weighted-average exercise price of $1.80 per share. During the three months ended September 30, 2024, 1,343 stock options were exercised and 64,961 stock options were forfeited at a weighted-average exercise price of $1.58 and $3.38 per share, respectively.Total compensation cost recognized for all stock option awards was as follows:â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Three Months Ended SeptemberÂ 30,Â â€‹Nine Months Ended SeptemberÂ 30,Â (in thousands)Â Â Â Â 2024Â Â Â Â 2023Â Â Â Â 2024Â Â Â Â 2023General and administrativeâ€‹$ 475â€‹$ 621â€‹$ 1,674â€‹$ 2,687Research and developmentâ€‹â€‹ 245â€‹â€‹ 303â€‹â€‹ 962â€‹â€‹ 1,427â€‹â€‹$ 720â€‹$ 924â€‹$ 2,636â€‹$ 4,114â€‹During the three months ended September 30, 2024, the Company transferred $0.1 million from the equity settled employee benefits reserve to accumulated deficit as a result of forfeited vested options. For these forfeited vested options there is no longer a requirement for a legal reserve as there are no limitations for distribution within equity.The fair value of the share options has been measured using the Black-Scholes model. The assumptions used in the measurement of the fair values and the weighted average of the share options granted during the nine months ended September 30, 2024:â€‹â€‹â€‹â€‹â€‹â€‹Â Â Â Â â€‹SeptemberÂ 30,Â 2024â€‹â€‹â€‹NL & USExpected annual average volatility Â Â Â Â â€‹92.6%Expected life, yearsÂ â€‹6.08Fair value of the share optionsâ€‹$1.22 - 1.57Exercise priceâ€‹$1.58 - 2.03Dividend yieldÂ â€‹ â€”Risk-free interest rateÂ â€‹4.07% - 4.34%Weighted average grant date fair valueÂ $1.2314Table of Contentsâ€‹The Company estimates volatility based on the historical volatility of its peer group. The unrecognized remaining stock-based compensation balance for shares issued from all option plans was approximately $2.4Â million as of September 30, 2024, which is expected to amortize over a weighted-average 0.79Â years.Note 9â€”Investments Our investments in debt securities consist of investments in U.S. Treasury securities, with maturities ranging from three to six months. All of these

investments are classified as held to maturity and recorded in current assets on our condensed consolidated interim statements of financial position at amortized cost. As of SeptemberÂ 30, 2024, the carrying value of our investments was $51.9 million, which approximates fair value. Given the high-quality ratings of these investments in debt securities, the Company has not recorded an allowance for credit losses as of September 30, 2024.Note 10â€“Subsequent Events In October 2024, the Company received notice from the RVO requiring a payment of $0.6 million within six weeks of the notice date. The remaining balance of $5.2 million, which is inclusive of principal and accrued interest, has been conditionally waived with a further decision to be made within one year from the notice date. Upon satisfaction of certain conditions related to pledged assets in the conditional waiver, the Company may request a permanent waiver from its remaining payment obligations to the RVO. The liability will remain classified as current, as the waiver lasts one year from the notice date.In October 2024, a milestone payment of $5.0 million from J&J was triggered under the terms of the J&J Agreement following the filing with health authorities to start a Phase 1 clinical trial. The $5.0 million is expected to be recorded as Revenue from contracts with customers.In December 2024, the Company decided to discontinue all cohorts of the LAVA-1207 clinical trial after determining that the potential signs of activity, including prostate-specific antigen reductions and several patients remaining on the study beyond six months, did not reach our internal benchmarks to continue. The LAVA-1207 clinical trial enrolled to dose level thirteen in the European Union and the United States and had no cytokine release syndrome events greater than grade 2 reported. An estimate of costs related to the discontinuance of the LAVA-1207 clinical trial cannot be made at this time. The Company has evaluated subsequent events through December 10, 2024. 15ExhibitÂ 99.2LAVA THERAPEUTICS, N.V.MANAGEMENTâ€™S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION ANDRESULTS OF OPERATIONSYou should read the following managementâ€™s discussion and analysis of our financial condition and results of operations in conjunction with our unaudited condensed consolidated interim financial statements, including the notes thereto, included with this report, as well as our audited consolidated financial statements as of, and for theÂ year ended, DecemberÂ 31, 2023, including the notes thereto, included in our annual report on FormÂ 20-F, filed with the Securities and Exchange Commission on March 20, 2024. The following discussion is based on our financial information prepared in accordance with International Accounting Standard 34, â€œInterim Financial Reporting.â€ Certain information and disclosures normally included in the consolidated financial statements prepared in accordance with International Financial Reporting Standards (IFRS) have been condensed or omitted. Throughout this managementâ€™s discussion and analysis, â€œwe,â€ us,â€ â€œour,â€ â€œLAVA,â€ and the â€œCompanyâ€ refer to LAVA Therapeutics N.V. and its consolidated subsidiaries, unless the context requires otherwise.Special NoteÂ Regarding Forward-Looking StatementsThis managementâ€™s discussion and analysis contains statements that constitute forward-looking statements. Many of the forward-looking statements contained in this managementâ€™s discussion and analysis can be identified by the use of forward-looking words such as â€œanticipate,â€ â€œbelieve,â€ â€œcould,â€ â€œexpect,â€ â€œshould,â€ â€œplan,â€ â€œintend,â€ â€œestimateâ€ and â€œpotential,â€ among others. Forward-looking statements appear in several places in this managementâ€™s discussion and analysis and include, but are not limited to, statements regarding our intent, belief or current expectations. Forward-looking statements are based on our managementâ€™s beliefs and assumptions and on information currently available to our management. Such statements are subject to risks and uncertainties, and actual results may differ materially from those expressed or implied in the forward-looking statements due to various important factors, including, but not limited to, those identified under the section titled â€œRisk Factorsâ€ in our annual report on Form 20-F. Forward-looking statements include but are not limited to, statements about:â—our operations as a biotechnology company with limited operating history and a history of operating losses;â—our plans to develop and commercialize our current and future product candidates;â—the initiation, timing, progress and results of our current and future preclinical studies and clinical trials and our research and development programs;â—our ability to successfully acquire or in-license additional product candidates on reasonable terms;â—our ability to maintain and establish collaborations or obtain additional funding;â—our ability to obtain regulatory approval of our current and future product candidates;â—our expectations regarding the potential market size and the rate and degree of market acceptance of our product candidates;â—our continued reliance on third parties to conduct clinical trials of our product candidate and future product candidates and manufacture our development candidates for preclinical studies and clinical trials;â—our ability to fund our working capital requirements and expectations regarding the sufficiency of our capital resources;â—the implementation of our business model and strategic plans for our business and product candidates;â—our ability to establish sales, marketing and distribution capabilities;â—our ability to enter into and maintain collaborations with third parties for the development or commercialization of our product candidates;â—our intellectual property position and the duration of our patent rights;â—our estimates regarding expenses, future revenues, capital requirements and our needs for additional financing;â—the impact of government laws and regulations on our business;â—our need to hire additional personnel and our ability to attract and retain such personnel;â—our ability to compete in the markets we serve;â—developments relating to our competitors and our industry; andâ—other risk factors discussed under â€œRisk Factorsâ€ in our annual report on Form 20-F.Forward-looking statements speak only as of the date they are made, and we do not undertake any obligation to update them in light of new information or future developments or to release publicly any revisions to these statements in order to reflect later events or circumstances or to reflect the occurrence of unanticipated events, except to the extent required by applicable law. In addition, there may be adverse effects on our business condition and results from rising interest rates, recent and potential future pandemics or other health crises, general economic and market conditions and overall fluctuations in the United States and international equity markets, including deteriorating market conditions due to investor concerns regarding inflation and international hostilities including the Russian invasion of Ukraine and the escalating conflict in the Middle East.OverviewWe are a clinical stage immuno-oncology company focused on developing our proprietary GammabodyÂ® platform of bispecific gamma delta (gd) T cell engagers to transform the treatment of cancer. Using our Gammabody platform, we are developing a portfolio of novel bispecific antibodies designed to engage and leverage the potency and precision of gd T cells to orchestrate a robust anti-tumor immune response and improve outcomes for cancer patients.We were incorporated in FebruaryÂ 2016 in the Netherlands and are currently headquartered in Utrecht, the Netherlands. In 2019, we established our wholly owned U.S. subsidiary, which began business in JanuaryÂ 2020. We established a wholly owned Australian subsidiary in August 2024. We have not generated any revenue from the sale of products. Since inception, we have incurred losses. As of September 30, 2024, we had an accumulated deficit of $168.9 million.We will transition from foreign private issuer to U.S. domestic filer status on January 1, 2025 and expect to incur increased costs associated with being a U.S. domestic filer, including expenses related to financial reporting, filing our annual report for the year ended December 31, 2024 on a Form 10-K, preparation of financial statements in accordance with U.S.Â Generally Accepted Accounting Principles, compliance with U.S. federal proxy rules, and additional resources and services we will require in order to comply with Nasdaq and SEC rules and requirements applicable to U.S. domestic filers.LAVA-1266LAVA-1266 is designed to target CD123+ tumor cells for the treatment of hematological malignancies, including acute myeloid leukemia (AML) and myelodysplastic syndrome (MDS). In October 2024, LAVA initiated a Phase 1 trial of LAVA-1266 in Australia. We plan to treat the first patient in this clinical trial shortly.LAVA-1207In 2022, we dosed the first patient in a first-in-human clinical trial evaluating LAVA-1207 in patients with metastatic castration-resistant prostate cancer (mCRPC). The open-label, multi-center, Phase 1/2a clinical trial was designed to evaluate safety, tolerability, pharmacokinetics, pharmacodynamics, immunogenicity and preliminary anti-tumor activity of LAVA-1207. The Phase 1 dose-escalation phase is designed to determine recommended Phase 2a dose(s) for optimization in Phase 2a. Once recommended Phase 2a dose(s) have been established, the trial was expected to expand into the Phase 2a portion to confirm safety and evaluate the preliminary anti-tumor activity of LAVA-1207 in patients with mCRPC.In February 2023, at the American Society of Clinical Oncology Genitourinary Cancers Symposium (ASCO GU), we reported clinical data for the ongoing Phase 1/2a clinical trial of LAVA-1207. For the first five cohorts, these initial data demonstrated predictable and linear pharmacokinetics and on-mechanism pharmacodynamics and a favorable safety profile. Preliminary signs of anti-tumor activity were observed at week 8, with iRECIST stable disease (iSD) in 8 out of 14 evaluable patients and PSA levels stabilizing or decreasing in several patients. iRECIST is the immune response evaluation criteria in solid tumors, a set of published rules that define whether tumors in cancer patients have improved, stayed the same or worsened during treatment. In June 2023, we introduced cohorts of patients who would receive one of two schedules of low-dose interleukin-2 (LDIL-2) beginning the day after LAVA-1207 dosing for the first four doses. LDIL-2 has the potential to increase the number of VÎ³9VÎ´2-T cells available for engagement by LAVA-1207. Three dose-limiting toxicities were reported in patients receiving LDIL-2 in addition to LAVA-1207 in cohort 7A2, a cohort with multiple doses of LDIL-2 per cycle but without step-dosing.Â Since introducing step-dosing, we have not observed any negative safety signals with the second dose of IL-2. â€‹In January 2024, we entered into a clinical trial collaboration and supply agreement with Merck & Co., Inc. to evaluate its anti-PD-1 therapy KEYTRUDAÂ® (pembrolizumab) in combination with LAVA-1207. Under the terms of this agreement, we have been provided with pembrolizumab for the dose escalation and expansion phases of LAVAâ€™s ongoing Phase 1/2a study of LAVA-1207 (NCT05369000) (KEYNOTE-F73). In the second quarter of 2024, we initiated dosing in the pembrolizumab combination arm and have treated our first patients. In December 2024, we decided to discontinue all cohorts of the LAVA-1207 clinical trial after determining that the potential signs of activity, including prostate-specific antigen reductions and several patients remaining on study beyond six months, did not reach our internal benchmarks to continue.Â The LAVA-1207 clinical trial enrolled to dose level thirteen (target dose of 45mg) in the European Union and the United States and had no cytokine release syndrome events greater than grade 2 reported. An estimate of costs related to the discontinuance of the LAVA-1207 clinical trial cannot be made at this time. PF-08046052 (formerly LAVA-1223)In 2022, we entered into an exclusive worldwide license agreement with Pfizer, Inc. (Pfizer Agreement) to develop, manufacture and commercialize PF-08046052, an advanced preclinical asset that utilizes our proprietary Gammabody technology to target epidermal growth factor receptor (EGFR)-expressing solid tumors. Under the terms of the Pfizer Agreement, we received a $50 million nonrefundable upfront payment in October 2022 and are eligible to receive up to approximately $650 million upon the achievement of development, regulatory and commercial milestones, as well as royalties ranging from the single digits to the mid-teens on future sales. The Pfizer Agreement also provided Pfizer with the opportunity to exclusively negotiate rights to apply our proprietary Gammabodyâ„¢ platform on up to two additional tumor targets, which Pfizer did not exercise. In 2023, we entered into a supply agreement with Pfizer to fulfill part of our obligations under the Pfizer Agreement and began shipping investigational drug supply to Pfizer in March 2023. As of September 30, 2023, all initial drug supply was shipped to Pfizer. In 2023, Pfizer received investigational new drug application clearance for PF-08046052 in advanced solid tumors from the FDA and initiated a Phase 1 trial (NTC0598133) of PF-08046052 to evaluate the safety and tolerability of this molecule as a monotherapy in advanced EGFR expressing solid tumors. In March 2024, Pfizer paid us $7 million for achieving a clinical development milestone.JNJ-89853413In 2020, we entered into a research collaboration and license agreement with J&J, for the discovery and development of novel bispecific antibody-based gamma delta T cell engagers for the treatment of cancer. In 2023, within the framework of the J&J Agreement, J&J selected a lead bispecific antibody utilizing the Gammabody platform for an undisclosed tumor associated antigen for development and we received a financial milestone payment. In the fourth quarter of 2024, J&J filed with health authorities to start a Phase 1 clinical trial, triggering a $5.0 million milestone payment to the Company. In December 2024, J&J will present a poster on JNJ-89853413 in a poster session at the American Society for Hematology Annual Meeting (ASH 2024)Comparison of the ThreeÂ Months Ended September 30, 2024 and 2023 (unaudited):Revenue from contracts with customers Our revenue from contracts with customers was zero and $0.1 million for the three months ended September 30, 2024 and 2023, respectively.In connection with the Pfizer Agreement, we recognized $0.1 million in revenue for the three months ended September 30, 2023, related to reimbursement for research activities and initial supply-related stability studies. Research and development expensesBelow were our research and development expenses:â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Â Â Â Three Months Ended Â Â Â Â Â Â Â Â â€‹â€‹â€‹SeptemberÂ 30,Â â€‹â€‹â€‹(in thousands)Â Â Â Â 2024Â Â Â Â 2023Â Â Â Â VariancePre-clinical and clinical trial expensesâ€‹$ 5,722â€‹$ 4,899â€‹$ 823Personnel-related expensesâ€‹Â 1,304â€‹Â 1,763â€‹Â (459)Research and development activities expensesâ€‹Â 678â€‹Â 325â€‹Â 353Facilities and other research and development expensesâ€‹Â 551â€‹Â 622â€‹Â (71)Share-based compensation expenseâ€‹Â 245â€‹Â 303â€‹Â (58)â€‹â€‹$ 8,500â€‹$ 7,912â€‹$ 588â€‹Research and development expenses were $8.5 million for the three months ended September 30, 2024, compared to $7.9 million for the three months ended September 30, 2023. Pre-clinical and clinical trial expenses increased by $0.8 million, primarily due to increased clinical trial activities for LAVA-1207 partly offset by reduced manufacturing costs for LAVA-1266 and other product candidates. Personnel-related expenses and non-cash share-based compensation expenses decreased by $0.5 million and $0.1 million, respectively, primarily due to research and development headcount reductions which occurred in the second half of 2023. Research and development activity expenses increased by $0.4 million, primarily due to increased patent costs and outsourced research costs. Facilities and other research and development expenses decreased by $0.1 million primarily due to reduced office and laboratory leases and related costs. General and administrative expensesBelow were our general and administrative expenses:â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Three Months Ended â€‹â€‹â€‹â€‹â€‹SeptemberÂ 30,Â â€‹â€‹â€‹(in thousands)Â Â Â Â 2024Â Â Â Â 2023Â Â Â Â VariancePersonnel-related expensesâ€‹$ 976â€‹$ 742â€‹$ 234Professional and consultant feesâ€‹Â 792â€‹Â 887â€‹Â (95)Insurance, facilities, fees and other related costsâ€‹â€‹ 542â€‹â€‹ 608â€‹â€‹ (66)Share-based compensation expenseâ€‹Â 475â€‹Â 621â€‹Â (146)â€‹â€‹$ 2,785â€‹$ 2,858â€‹$ (73)â€‹General and administrative expenses were $2.8 million for the three months ended September 30, 2024, compared to $2.9 million for the three months ended September 30, 2023. Personnel-related expenses increased by $0.2 million. Professional and consultant fees decreased by $0.1 million. Insurance, facilities, fees and other related costs decreased by $0.1 million, primarily due to reduced directors and officers insurance premiums and reduced office lease costs. Non-cash share-based compensation expenses decreased by $0.1 million. Interest income, netInterest income, net was comparable at $0.8 million for the three months ended September 30, 2024, and 2023. Interest income, net includes interest income from investments, net of interest on borrowings associated with our RVO Innovation Credit and lease interest expense.Foreign currency exchange (loss) gain, netFor the three months ended September 30, 2024 and 2023, foreign currency exchange loss increased by $2.8 million, from a gain of $1.1 million during the three months ended September 30, 2023 to a loss of $1.7 million during the three months ended September 30, 2024. This loss was due to the impact of the fluctuation of the USD currency rate compared to the Euro on transaction gains and losses on cash and investments and other transactions denominated in USD held and occurring in the Euro functional currency entity.Comparison of the NineÂ Months Ended September 30, 2024 and 2023 (unaudited):Revenue from contracts with customers Our revenue from contracts with customers was $7.0 million and $6.4 million for the nine months ended September 30, 2024 and 2023, respectively. In connection with the Pfizer Agreement, we recognized $7.0 million in revenue for the nine months ended September 30, 2024, primarily related to the achievement by Pfizer of a clinical development milestone for PF-08046052.In connection with the Pfizer Agreement, we recognized $3.9 million in revenue for the nine months ended September 30, 2023, related to reimbursement for research activities and delivery of initial supply. In connection with the J&J Agreement, we recognized $2.5 million in revenue for the nine months ended September 30, 2023, related to a triggered milestone payment.Cost of providing services and sales of goodsâ€‹Our cost of providing services and sales of goods was zero and $3.3 million for the nine months ended September 30, 2024 and 2023, respectively. The $3.3 million for the nine months ended September 30, 2023 related to the cost of the initial supply delivery and related stability studies. Of this amount $0.8 million related to costs of services related to stability studies for the product materials and $2.5 million related to costs of product materials.â€‹Research and development expensesBelow were our research and development expenses:â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹For the Nine Months Ended â€‹â€‹â€‹â€‹â€‹SeptemberÂ 30,Â â€‹â€‹â€‹(in thousands)Â Â Â Â 2024Â Â Â Â 2023Â Â Â Â VariancePre-clinical and clinical trial expensesâ€‹$ 12,836â€‹$ 20,542â€‹$ (7,706)Personnel-related expensesâ€‹Â 3,830Â Â 5,346â€‹Â (1,516)Research and development activities expensesâ€‹Â 2,024Â Â 1,297â€‹Â 727Facilities and other research and development expensesâ€‹Â 1,192Â Â 1,842â€‹Â (650)Share-based compensation expenseâ€‹Â 962Â Â 1,427â€‹Â (465)â€‹â€‹$ 20,844â€‹$ 30,454â€‹$ (9,610)â€‹Research and development expenses were $20.8 million for the nine months ended September 30, 2024, compared to $30.5 million for the nine months ended September 30, 2023. Pre-clinical and clinical trial expenses decreased by $7.7 million, primarily due to reduced manufacturing scale-up costs and reduced activities of the clinical trials as a result of the discontinuation of LAVA-051, announced in September 2023. Personnel-related expenses and non-cash share-based compensation expenses decreased by $1.5 million and $0.5 million, respectively, primarily due to research and development headcount reductions which occurred in the second half of 2023. Research and development activity expenses increased by $0.7 million, primarily due to increased patent costs. Facilities and other research and development expenses decreased by $0.6 million primarily due to reduced office and laboratory leases and related costs. General and administrative expensesBelow were our general and administrative expenses:â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹For the Nine Months Ended â€‹â€‹â€‹â€‹â€‹SeptemberÂ 30,Â â€‹â€‹â€‹(in thousands)Â Â Â Â 2024Â Â Â Â 2023Â Â Â Â VariancePersonnel-related expensesâ€‹$ 2,937â€‹$ 2,932â€‹$ 5Professional and consultant feesâ€‹Â 2,356â€‹Â 2,667â€‹Â (311)Insurance, facilities, fees and other related costsâ€‹Â 1,778â€‹Â 2,159â€‹Â (381)Share-based compensation expenseâ€‹â€‹ 1,674â€‹â€‹ 2,687â€‹â€‹ (1,013)â€‹â€‹$ 8,745â€‹$ 10,445â€‹$ (1,700)â€‹General and administrative expenses were $8.7 million for the nine months ended September 30, 2024, compared to $10.4 million for the nine months ended September 30, 2023.

Professional and consultant fees decreased by $0.3 million, primarily due to lower legal fees partly offset by increased audit and compliance fees. Insurance, facilities, fees and other related costs decreased by $0.4 million, primarily due to reduced directors and officers insurance premiums and reduced office lease costs. Non-cash share-based compensation expenses decreased by $1.0 million, primarily due to a reduction in general and administrative headcount, which occurred in the second half of 2023. Interest income, netInterest income, net was $2.4 million for the nine months ended September 30, 2024, compared to $2.1 million for the nine months ended September 30, 2023. The increase in interest income was primarily due to higher interest yields on our investments in 2024. Interest income, net includes interest income from investments, net of interest on borrowings associated with our RVO Innovation Credit and lease interest.Foreign currency exchange (loss) gain, netFor the nine months ended September 30, 2024 and 2023, foreign currency exchange loss, net increased by $1.1 million, from a gain of $0.4 million during the nine months ended September 30, 2023 to a loss of $0.7 million during the nine months ended September 30, 2024. This increase was due to the impact of the fluctuation of the USD currency rate compared to the Euro on transaction gains and losses on cash and investments and other transactions denominated in USD held and occurring in the Euro functional currency entity.Liquidity and Capital ResourcesAs of September 30, 2024, we had cash, cash equivalents and investments totaling $78.9 million, compared to cash, cash equivalents and investments of $95.6 million as of DecemberÂ 31, 2023. We have historically funded our operations primarily through the issuance of preference shares prior to our IPO and from the sale of common shares in our IPO in March 2021, and proceeds from the Pfizer Agreement and J&J Agreement. Our expenditures are primarily related to research and development activities and general and administrative activities to support business operations.In April 2022, we entered into an Equity Distribution Agreement (EDA) with JMP Securities LLC (JMP) under which JMP, as our exclusive agent, at our discretion and at such times that we may determine from time to time, may sell over a three-year period from the execution of the agreement up to a maximum of $50 million of shares of our common stock. We have not sold any of our common shares under the EDA to date.In September 2022, we entered into the Pfizer Agreement for the development, manufacture and commercialization of PF-08046052 (formerly LAVA-1223), an advanced preclinical asset that utilizes LAVAâ€™s proprietary Gammabody technology to target EGFR-expressing solid tumors. Under the terms of the agreement, we received a $50 million nonrefundable upfront payment in October 2022. In March 2024, we received a milestone payment of $7.0 million from Pfizer following the achievement of a clinical milestone for PF-08046052.In October 2024, a milestone payment of $5.0 million from J&J was triggered under the terms of the J&J Agreement following the confirmation of its filing with health authorities to start a Phase 1 clinical trial. Cash and cash equivalents, and short-term marketable securities are financial instruments that potentially subject the Company to concentrations of credit risk. As of September 30, 2024 cash consisted of cash deposited with four financial institutions, with account balances exceeding federally insured limits with three institutions. As of December 31, 2023, cash consisted of cash deposited with three financial institutions; account balances exceeded federally insured limits. â€‹Based on our current operating plan, we believe that our existing cash, cash equivalents and investments as of September 30, 2024 are sufficient to meet our projected cash requirements for at least 12Â months from the date of this report. However, our operating plan may change as a result of many factors currently unknown to us, and we may need to seek additional funds sooner than planned. Our future funding requirements will depend on many factors, including, but not limited to, our ability to:â—continue the ongoing and planned development of our product candidates, including LAVA-1266;â—initiate, conduct and complete any ongoing, anticipated or future preclinical studies and clinical trials for our current and future product candidates;â—develop processes and scale manufacturing production for our current and future product candidates in accordance with cGMP;â—seek regulatory and marketing approvals for LAVA-1266 and any of our other development candidates that successfully complete clinical trials;â—discover and develop additional bispecific gd engagers and make further investments in our Gammabody platform to identify additional product candidates;â—maintain, protect and expand our intellectual property portfolio; including costs associated with opposing and invalidating competitor patents and licensing other technologies for our product candidates;â—establish a sales, marketing, manufacturing and distribution, supply chain and other commercial infrastructure in the future to commercialize any current or future product candidate for which we may obtain marketing approval;â—expand our operations in the United States, Europe and Australia;â—add clinical, scientific, operational, financial and management information systems and personnel, including personnel to support our product development and planned future commercialization efforts;â—acquire or in-license additional product candidates and technologies;â—develop a potential companion diagnostic;â—incur additional legal, accounting and other expenses associated with the transition from foreign private issuer to U.S. domestic filer status; â—address any events outside of our control, including, but not limited to, outbreaks of infectious diseases; and â—face general economic and market conditions and overall fluctuations in the United States and international equity markets, such as deteriorating conditions due to investor concerns regarding inflation and the Russian invasion of Ukraine, the escalating conflict in the Middle East, and other geopolitical conditions. Additional funds may not be available when we need them on terms that are acceptable to us, or at all. If adequate funds are not available to us on a timely basis, we may be required to delay, limit, scale back or cease our research and development activities, preclinical studies and clinical trials for our product candidates and our establishment and maintenance of sales and marketing capabilities or other activities that may be necessary to commercialize our product candidates.The following is a summary of our cash flows:â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹For the Nine Months Ended â€‹â€‹SeptemberÂ 30,Â (in thousands)Â Â Â Â 2024Â Â Â Â 2023â€‹Net cash used in operating activitiesâ€‹$ (18,139)â€‹$ (28,507)â€‹Net cash provided by (used in) investing activitiesâ€‹Â 1,504â€‹Â (8,948)â€‹Net cash used in financing activitiesâ€‹Â (363)â€‹Â (736)â€‹Net decrease in cash and cash equivalentsâ€‹$ (16,998)â€‹$ (38,191)â€‹â€‹Cash Flows Used in Operating ActivitiesNet cash used in operating activities for the nineÂ months ended September 30, 2024, was $18.1 million compared to $28.5 million for the nine months ended September 30, 2023. During the nine months ended September 30, 2024, we incurred net losses of $20.9 million and had $2.0 million amortization of premium on investments, primarily offset by noncash share-based compensation expenses of $2.6 million, foreign currency loss of $0.7 million, and changes in working capital of $0.6 million. During the nine months ended September 30, 2023, we incurred net losses of $35.3 million and had $1.2 million amortization of premium on investments, primarily offset by noncash share-based compensation expenses of $4.1 million and changes in working capital of $3.2 million. The reduction in net losses in the nine months ended September 30, 2024, compared to 2023, was largely due to the receipt of the clinical milestone of $7.0 million from Pfizer in 2024, reduced research and development expenses as a result of the discontinuation of LAVA-051, announced in September 2023, and reduced general and administrative expenses.Cash Flows Provided by (Used in) Investing ActivitiesCash flows provided by investing activities for the nine months ended September 30, 2024, were $1.5 million and primarily consisted of $77.0 million of investments matured, offset by $75.6 million purchases of investments. Cash flows used in investing activities for the nine months ended September 30, 2023 were $8.9 million and primarily consisted of $53.9 million of purchase of investments and $0.7 million of equipment purchases, offset by $45.7 million of investments matured. Cash Flows Used in Financing ActivitiesCash flows used in financing activities for the nineÂ months ended September 30, 2024, were $0.4 million and primarily consisted of $0.4 million in principal payments on operating lease liabilities. Cash flows used in financing activities for the nine months ended September 30, 2023, were $0.7 million and consisted of $0.7 million in principal payments on operating lease liabilities. Off-Balance Sheet ArrangementsWe have not entered into any off-balance sheet arrangements or any holdings in variable interest entities.Quantitative and Qualitative Disclosures about Market RiskWe are exposed to a variety of risks in the ordinary course of our business, including, but not limited to, foreign currency risk, interest rate risk, credit risk and liquidity risk. We regularly assess each of these risks to minimize any adverse effects on our business as a result of those factors.Foreign Currency RiskWe are exposed to foreign exchange risk arising from various currency exposures, primarily with respect to USD. We have received payments in USD under our collaborations, and the proceeds from our initial public offering in MarchÂ 2021 were in USD and we execute some transactions in USD. As a result, we are exposed to volatility in the condensed consolidated interim statements of profit and loss related to USD amounts and transactions occurring in a Euro-functional entity, the impacts of which we have disclosed above. We regularly assess our foreign currency risk, maintain cash positions in the currencies in which we expect to incur the majority of our future expenses and may engage in hedging activities consistent with our investment policy to minimize this risk and preserve our capital.Interest Rate RiskWe have interest-bearing debt with third parties. In addition, while we have no derivatives or financial assets and liabilities measured at fair value, our exposure to interest rate risk primarily relates to the interest rates for our positions of cash and cash equivalents, including short-term marketable securities. Our future interest income from interest-bearing bank deposits and short-term investments may fall short of expectations due to changes in interest rates. We do not consider the effects of interest rate fluctuations to be a material risk to our financial position.We have adopted an investment policy with the primary purpose of preserving capital, fulfilling our liquidity needs and diversifying the risks associated with cash and marketable securities. This investment policy establishes minimum ratings for institutions with which we hold cash, cash equivalents and marketable securities, as well as rating and concentration limits for marketable securities that we may hold.Credit RiskWe consider all of our material counterparties to be creditworthy. While the concentration of credit risk may be significant, we consider the credit risk for each of our counterparts to be low. Our exposure to credit risk primarily relates to our cash and cash equivalents, comprising bank deposits and short-term marketable securities with a maturity of sixÂ months or less at the date of acquisition. The credit risk on bank deposits is limited because the counterparties holding significant deposits are banks with high credit ratings assigned by international credit rating agencies. Our banks are reviewed on a regular basis, and our deposits may be transferred during theÂ year to mitigate credit risk. We have considered the risk of expected credit loss on our cash deposits, including the hypothetical impact arising from the probability of default, considering in conjunction with the expected loss given default from banks with similar credit ratings and attributes. In line with previous periods, our assessment did not reveal a material impairment loss, and accordingly, no provision for expected credit loss has been made. We hold a portion of our bank deposits in a money market fund invested in short-term U.S. Treasury securities to further diversify the credit risk. For other financial assets, including deposits and receivables, we consider the credit risk to be low and no provision for expected credit loss has been made.Liquidity RiskWe manage our liquidity risk by maintaining adequate cash reserves and banking facilities, continuously monitoring our cash forecasts and actual cash flows and matching the maturity profiles of financial assets and liabilities. We monitor the risk of a shortage of funds using a liquidity planning tool to ensure enough funds are available to settle liabilities as they fall due.Historically we have addressed the risk of insufficient funds through the proceeds from our SeriesÂ C financing and our IPO in MarchÂ 2021, and research and license agreements with strategic partners.Critical Accounting EstimatesThe preparation of financial statements in conformity with IFRS requires the use of certain critical accounting estimates and requires management to exercise its judgment in the process of applying our accounting policies. The areas involving a higher degree of judgment or complexity or areas where assumptions and estimates are significant to the unaudited condensed consolidated interim financial statements are disclosed in NoteÂ 3 to our condensed consolidated interim financial statements. The interim financial data as of September 30, 2024 and 2023 are unaudited. In the opinion of management, the interim financial data includes all adjustments, consisting only of normal recurring adjustments, necessary to a fair statement of the results for the interim periods.â€‹RISK FACTORSThe risk factors set forth under the captionÂ â€œRisk Factorsâ€Â in Item 3 of our annual report on Form 20-F filed by the Company on March 20, 2024 shall be deemed to be incorporated by reference herein and to be a part hereof from the date on which this report is filed, to the extent not superseded by documents or reports subsequently filed or furnished. Additional risks and uncertainties not currently known to the Company, or that the Company currently deems to be immaterial, may also affect its business, financial condition and/or future operating results.â€‹Exhibit 99.3â€‹â€‹LAVA Reports Third Quarter 2024 Financial Results and Announces Pipeline Reprioritization and Cash Runway Extension into 2027â€‹â—Reprioritized pipeline to focus on LAVA-1266, with continued support for partnered programs with Pfizer (PF-08046052) and Johnson and Johnson (JNJ-89853413) and plan to discontinue development of LAVA-1207 â—Fourth quarter pipeline advances include initiation of the Phase 1 trial for LAVA-1266, for hematologic malignancies, and a $5.0 million milestone payment from Johnson and Johnsonâ—Cash runway extended into 2027, based on a cash balance of $78.9 million, as of September 30, 2024 Utrecht, The Netherlands, and Philadelphia, PA, US â€“ December 10, 2024 â€“ LAVA Therapeutics N.V. (NASDAQ: LVTX, â€œLAVA,â€ â€œthe Companyâ€), a clinical-stage immuno-oncology company focused on developing its proprietary GammabodyÂ® platform of bispecific gamma delta T cell engagers, reported financial results for the third quarter ended September 30, 2024 and announced a strategic pipeline reprioritization.â€œLAVAâ€™s goal is to develop immuno-oncology medicines to improve the lives of cancer patients. While we are disappointed that LAVA-1207 did not reach our predetermined success criteria, we are reprioritizing our pipeline to focus on LAVA-1266, for acute myeloid leukemia (AML) and myelodysplastic syndrome (MDS) and will continue to support our partnered programs. We are pleased by the progress of our partnered programs, including a preclinical data presentation by Johnson and Johnson at ASH 2024 and ongoing enrollment in the Phase 1 program by Pfizer,â€ said Stephen Hurly, President and Chief Executive Officer of LAVA Therapeutics. â€œLAVA is well capitalized, with approximately $79 million in cash, and with this pipeline reprioritization, we expect our cash balance to fund the Company into 2027.â€ We thank the patients, investigators, and our employees for supporting the LAVA-1207 clinical study,â€ said Charlie Morris, MD, Chief Medical Officer of LAVA Therapeutics. â€œThe longer time to progression, with several patients on trial beyond 6 months, and duration of treatment observed for patients with higher circulating gamma delta2 T cells is consistent with the mechanism of action and supports continued clinical investigation of the platform.â€Portfolio Highlights: LAVA-1266 â€“ In Phase 1 Trial (ACTRN12624001214527)Designed to target CD123+ tumor cells for the treatment of hematological malignanciesâ—Key indications: Acute myeloid leukemia (AML) and myelodysplastic syndrome (MDS)â—Current Status: Phase 1 dose escalation study initiated in Australia Johnson & Johnson Partnered Program (JNJ-89853413) â€“ Phase 1 Trial (NCT06618001)Designed to target CD33 and VÎ´2 T cells with a bispecific gamma delta T cell engager â—Key Indications: include hematological cancersâ—Current Status: Johnson and Johnson has filed with health authorities to start in a Phase 1 study. Johnson & Johnson presented preclinical data for JNJ-89853413 at the Annual Meeting of the American Society of Hematology (ASH 2024) on December 7, 2024 (Abstract 2054: 2054). â—Milestone: Development milestone of $5 million received from Johnson and Johnson in Q4 2024 related to the IND filing for JNJ-89853413Pfizer Partnered Program â€“(PF08046052) â€“ In Phase 1 Trial (NCT05983133)Â Potential first-in-class epidermal growth factor receptor (EGFR) and bispecific gamma delta T cell receptor-targeted therapy for solid tumors â—Key Indications: Include colorectal cancer (CRC), non-small cell lung cancer (NSCLC), head and neck squamous cell carcinoma (HNSCC) and pancreatic ductal adenocarcinoma (PDAC)â—Current Status: Monotherapy Phase 1 dose escalation study underway to evaluate the safety and tolerability in advanced EGFR-expressing solid tumorsâ—Milestone: Clinical development milestone of $7 million received from Pfizer in Q1 2024LAVA-1207 â€“ Discontinued Designed to target prostate-specific membrane antigen (PSMA)-positive metastatic castration-resistant prostate cancer (mCRPC) â—The Phase 1 study of LAVA-1207 did not reach LAVAâ€™s internal benchmarks and the program and drug will be made available for patients currently receiving LAVA-1207 for as long as considered necessary by their treating physician. The decision to discontinue LAVA-1207 is not due to safety concerns. Clinical signals recorded in several patients, including PSA reductions and extended time on study for patients with higher baseline circulating Vd2 T cells as well as the encouraging safety and tolerability profile are consistent with the intended mechanism of action and support further investigation of the platform. Learnings, especially related to the favorable overall safety profile and tolerability of LAVA-1207, will be incorporated into further development of the Companyâ€™s pipeline programs.Third Quarter 2024 Financial Results â—As of September 30, 2024, LAVA had cash, cash equivalents and investments totaling $78.9 million, compared to cash, cash equivalents and investments of $95.6 million as of DecemberÂ 31, 2023. The Company believes its current cash, cash equivalents and investments will be sufficient to fund operations into 2027. â—Revenue from contracts with customers was zero and $0.1 million for the quarters ended September 30, 2024 and 2023, respectively, and $7.0 million and $6.4 million for the nine months ended September 30, 2024 and 2023, respectively. Revenue of $0.1 million for the quarter ended September 30, 2023 was related to reimbursement for research activities and initial stability studies for clinical supplies. â—Research and development expenses were $8.5 million and $7.9 million for the quarters ended September 30, 2024, and 2023, respectively, and $20.8 million and $30.5 million for the nine months ended September 30, 2024 and 2023, respectively. The increase for the quarter ended September 30, 2024, as compared to 2023 was the result of increased pre-clinical and clinical trial expenses due to increased clinical trial activities for LAVA-1207, partly offset by reduced manufacturing costs for LAVA-1266 and other product candidates. The decrease between the nine months ended September 30, 2024 and 2023, respectively, was primarily due to lower pre-clinical and clinical trial expenses due to the discontinuation of LAVA-051, announced in June 2023, and reduced personnel-related expenses due to a reduction in research and development headcount in the second half of 2023.â—General and administrative expenses were $2.8 million and $2.9 million for the quarters ended September 30, 2024 and 2023, respectively, and $8.7 million and $10.4 million for the nine months ended September 30, 2024 and 2023, respectively. The decrease in both periods was primarily due to lower non-cash share-based compensation expenses and personnel-related expenses due to a reduction in general and administrative headcount in the second half of 2023. â—Net losses were $12.3 million and $8.8 million, or $0.46 and $0.34 net loss per share, for the quarters ended September 30, 2024 and 2023, respectively, and $21.1 million and $35.5 million, or $0.79 and $1.35 net loss per share, for the nine months ended September 30, 2024 and 2023, respectively.LAVA will transition from foreign private issuer to U.S. domestic filer status beginning on January 1, 2025. The Company

expects to incur increased costs associated with this transition, including expenses related to financial reporting, preparation of financial statements in accordance with U.S.Â GAAP, and compliance with U.S. federal proxy rules.â€‹LAVA Therapeutics N.V.Condensed Consolidated Interim Statements Lossand Comprehensive Loss(in thousands, except share and per share amounts) (unaudited)â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Â Â Â Â Â Â Â Â Â Â Â Â Three Months Ended Â Â Â Â Nine Months Ended â€‹â€‹â€‹â€‹â€‹SeptemberÂ 30,Â â€‹SeptemberÂ 30,Â â€‹â€‹Â Â Â Â NotesÂ Â Â Â 2024Â Â Â Â 2023Â Â Â Â 2024Â Â Â Â 2023â€‹Revenue:Â Â Â Â â€‹Â Â Â â€‹Â Â Â â€‹Â Â Â â€‹Â Â â€‹Revenu from contracts with customersÂ 5â€‹$ â€”â€‹$ 53â€‹$ 6,992â€‹$ 6,416â€‹Cost of sales of goodsâ€‹5â€‹â€‹ â€”â€‹â€‹ â€”â€‹â€‹ â€”â€‹â€‹ (2,546)â€‹Cost of providing servicesâ€‹5â€‹â€‹ â€”â€‹â€‹ (10)â€‹â€‹ â€”â€‹â€‹ (782)â€‹Gross profitÂ Â Â â€‹Â â€”â€‹Â 43â€‹Â 6,992â€‹Â 3,088â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Operating expenses:Â Â Â â€‹Â Â Â â€‹Â Â Â â€‹Â Â Â â€‹Â Â Â â€‹Research and developmentÂ 6â€‹Â (8,500)â€‹Â (7,912)â€‹Â (20,844)â€‹Â (30,454)â€‹General and administrativeÂ 7â€‹Â (2,785)â€‹Â (2,858)â€‹Â (8,745)â€‹Â (10,445)â€‹Total operating expensesÂ Â Â â€‹Â (11,285)â€‹Â (10,770)â€‹Â (29,589)â€‹Â (40,899)â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Operating lossÂ Â Â â€‹Â (11,285)â€‹Â (10,727)â€‹Â (22,597)â€‹Â (37,811)â€‹Interest income, netÂ â€‹â€‹Â 805â€‹Â 809â€‹Â 2,426â€‹Â 2,124â€‹Foreign currency exchange (loss) gain, netÂ â€‹â€‹Â (1,720)â€‹Â 1,132â€‹Â (723)â€‹Â 429â€‹Total non-operating (loss) incomeÂ Â Â â€‹Â (915)â€‹Â 1,941â€‹Â 1,703â€‹Â 2,553â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Loss before income taxÂ Â Â â€‹Â (12,200)â€‹Â (8,786)â€‹Â (20,894)â€‹Â (35,258)â€‹Income tax expenseÂ â€‹â€‹Â (96)â€‹Â (50)â€‹Â (250)â€‹Â (218)â€‹Loss for the periodÂ Â Â â€‹$ (12,296)â€‹$ (8,836)â€‹$ (21,144)â€‹$ (35,476)â€‹Items that may be reclassified to profit or lossâ€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Foreign currency translation adjustmentÂ â€‹â€‹Â 1,688â€‹Â (1,385)â€‹Â 296â€‹Â (82)â€‹Total comprehensive lossÂ Â Â â€‹$ (10,608)â€‹$ (10,221)â€‹$ (20,848)â€‹$ (35,558)â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹Â Â â€‹â€‹â€‹â€‹â€‹â€‹â€‹Loss per share:Â Â Â â€‹Â Â Â â€‹Â â€‹â€‹Â Â Â â€‹Â Â Â â€‹Loss per share, basic and dilutedâ€‹â€‹â€‹$ (0.46)â€‹$ (0.34)â€‹$ (0.79)â€‹$ (1.35)â€‹Weighted-average common shares outstanding, basic and dilutedâ€‹â€‹â€‹Â 26,846,006â€‹Â 26,289,087â€‹Â 26,814,113â€‹Â 26,289,087â€‹â€‹â€‹â€‹LAVA Therapeutics N.V.Condensed Consolidated Statements of Financial Position(in thousands) (unaudited)â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹â€‹SeptemberÂ 30,Â â€‹DecemberÂ 31,Â â€‹Â Â Â Â NotesÂ Â Â Â 2024Â Â Â Â 2023AssetsÂ Â Â Â â€‹Â Â Â â€‹Â Â Non-current assets:Â Â Â Â â€‹Â Â Â â€‹Â Â Property and equipment, netÂ â€‹â€‹$ 1,188â€‹$ 1,602Right-of-use assetsÂ â€‹â€‹Â 517â€‹Â 892Deferred tax assetsÂ Â Â â€‹Â â€”â€‹Â â€”Other non-current assets and security depositsÂ Â Â â€‹Â 114â€‹Â 319Total non-current assetsÂ Â Â â€‹Â 1,819â€‹Â 2,813Current assets:Â Â Â â€‹Â Â Â â€‹Â Â Â Receivables and otherÂ â€‹â€‹Â 536â€‹Â 1,459Prepaid expenses and other current assetsÂ â€‹â€‹Â 1,629â€‹Â 1,627VAT receivableÂ â€‹â€‹Â 489â€‹Â 240Investmentsâ€‹9â€‹â€‹ 51,921â€‹â€‹ 51,340Cash and cash equivalentsÂ â€‹â€‹Â 26,963â€‹Â 44,231Total current assetsÂ Â Â â€‹Â 81,538â€‹Â 98,897Total assetsÂ Â Â â€‹$ 83,357â€‹$ 101,710Equity and LiabilitiesÂ Â Â â€‹Â Â Â â€‹Â Â Â Equity:Â Â Â â€‹Â Â Â â€‹Â Â Â Share capitalÂ â€‹â€‹$ 3,716â€‹$ 3,715Equity-settled employee benefits reserveÂ 8â€‹Â 14,360â€‹Â 12,005Foreign currency translation reserveÂ â€‹â€‹Â (10,603)â€‹Â (10,899)Additional paid-in capitalÂ â€‹â€‹Â 194,450â€‹Â 194,424Accumulated deficitÂ Â Â â€‹Â (168,930)â€‹Â (148,067)Total equityÂ Â Â â€‹Â 32,993â€‹Â 51,178Non-current liabilities:Â Â Â â€‹Â Â Â â€‹Â Â Â Deferred revenueÂ 5â€‹Â 35,000â€‹Â 35,000Lease liabilitiesÂ â€‹â€‹Â 164â€‹Â 591Total non-current liabilitiesÂ Â Â â€‹Â 35,164â€‹Â 35,591Current liabilities:Â Â Â â€‹Â Â Â â€‹Â Â Â Trade payables and otherÂ â€‹â€‹Â 2,815â€‹Â 4,446Borrowingsâ€‹â€‹â€‹â€‹ 5,756â€‹â€‹ 5,282Lease liabilitiesÂ â€‹â€‹Â 370â€‹Â 440Accrued expenses and other current liabilitiesÂ â€‹â€‹Â 6,259â€‹Â 4,773Total current liabilitiesÂ Â Â â€‹Â 15,200â€‹Â 14,941Total liabilitiesÂ Â Â â€‹Â 50,364â€‹Â 50,532Total equity and liabilitiesÂ Â Â â€‹$ 83,357â€‹$ 101,710â€‹â€‹â€‹â€‹About LAVA Therapeutics LAVA Therapeutics N.V. is a clinical-stage immuno-oncology company focused on advancing its proprietary GammabodyÂ® platform to develop a portfolio of bispecific gamma-delta T cell engagers for the potential treatment of solid tumors and hematologic malignancies. The Company utilizes bispecific antibodies engineered to selectively kill cancer cells by triggering VÎ³9VÎ´2 (Vgamma9 Vdelta2) T cell anti-tumor effector functions upon cross-linking to tumor-associated antigens.LAVAâ€™s pipeline includes three internal and partnered clinical stage bispecific gamma-delta T cell engagers for the treatment of solid tumor and hematological cancers including LAVA 1266, targeting CD123+ cancers; PF-08046052, targeting EGFR (NCT05983133); and JNJ-89853413, targeting hematological cancers (NCT06618001). The pipeline also includes pre-clinical programs. For more information on LAVA, please visit our website at www.lavatherapeutics.com, or follow us on LinkedIn, X, and YouTube.GammabodyÂ® is a registered trademark of LAVA Therapeutics N.V.LAVAâ€™s Cautionary Note on Forward-Looking Statements This press release contains forward-looking statements, regarding the Companyâ€™s business and clinical development plans including the timing and results of clinical trials. Words such as â€œanticipateâ€, â€œbelieveâ€, â€œcouldâ€, â€œwillâ€, â€œmayâ€, â€œexpectâ€, â€œshouldâ€, â€œplanâ€, â€œintendâ€, â€œestimateâ€, â€œpotentialâ€, â€œsuggestsâ€, and similar expressions (as well as other words or expressions referencing future events, conditions or circumstances) are intended to identify forward-looking statements. These forward-looking statements are based on LAVAâ€™s expectations and assumptions as of the date of this press release and are subject to various risks and uncertainties that may cause actual results to differ materially from these forward-looking statements. Forward-looking statements contained in this press release include but are not limited to statements relating to the therapeutic potential, development strategy and potential uses of LAVAâ€™s product candidates including the timing of initiation of clinical trials and achievement of clinical milestones, and the Companyâ€™s ability to realize the expected benefits of its strategic pipeline reprioritization, including generation of clinical data for LAVA-1266, LAVAâ€™s cash runway and the sufficiency of resources to pursue development activities, expectations related to increased costs associated with transitioning from foreign private issuer status to a U.S. domestic filer status, availability of information regarding clinical development plans, progress and data from clinical trials, and the ability of LAVAâ€™s product candidates to treat various tumor targets and improve patient outcomes. Many factors, risks and uncertainties may cause differences between current expectations and actual results, including, among other things, the Companyâ€™s ability to leverage its initial programs to develop additional product candidates using its GammabodyÂ® platform, and the failure of LAVAâ€™s collaborators to support or advance collaborations or LAVAâ€™s product candidates, the timing and results of LAVAâ€™s research and development programs and preclinical and clinical trials, the possibility that clinical trials may fail to establish sufficient efficacy, the risk that adverse events or safety signals may occur in clinical trials, the risk that results obtained in clinical trials to date may not be indicative of results obtained in ongoing or future trials, the risk that adverse regulatory actions or other setbacks could occur in clinical trials even after promising results in earlier clinical trials or preclinical studies, the Companyâ€™s ability to obtain regulatory approval for and commercialize its product candidates, and the risk that setbacks in development could occur as a result of the difficulty and uncertainty of pharmaceutical product development and other factors. There may be adverse effects on the Companyâ€™s business condition and results from general economic and market conditions and overall fluctuations in the United States and international equity markets, including as a result of inflation, heightened interest rates, recent and potential future pandemics and other health crises, and hostilities, including between Russia and escalating tension in the Middle East. These and other risks are described in greater detail under the caption â€œRisk Factorsâ€ in LAVAâ€™s most recent Annual Report on Form 20-F and other filings the Company makes with the Securities and Exchange Commission. LAVA assumes no obligation to update any forward-looking statements contained herein to reflect any change in expectations, even as new information becomes available.â€‹CONTACTSInvestor Relationsir@lavatherapeutics.com â€‹LifeSci Advisors (IR/Media)Joyce AllaireJallaire@lifesciadvisors.com