UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 6-K

REPORT OF FOREIGN PRIVATE ISSUER
PURSUANT TO RULE 13a-16 OR 15d-16
UNDER THE SECURITIES EXCHANGE ACT OF 1934

Date as December 10, 2024

Commission File Number 001-35428

IMMUTEP LIMITED
(Exact Name as Specified in its Charter)

N/A
(Translation of Registrant's Name)

Level 32, Australia Square
264 George Street, Sydney
NSW 2000, Australia
(Address of principal executive office)

Indicate by check mark whether the registrant files or will file annual reports under cover of Form 20-F or Form 40-F.

Form 20-F ☒ Form 40-F ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1): ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7): ☐

Indicate by check mark whether by furnishing the information contained in this Form, the registrant is also thereby furnishing the information to the Commission pursuant to Rule 12g3-2(b) under the Securities Exchange Act of 1934.

Yes ☐ No ☒

If "Yes" is marked, indicated below the file number assigned to the registrant in connection with Rule 12g3-2(b): Not applicable.

EXHIBIT INDEX

Exhibit	Description of Exhibit
99.1	Immutep Announces Initiation of TACTI-004 Phase III Trial in First Line Non-Small Cell Lung Cancer

SIGNATURES

Pursuant 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.

Date: December 10, 2024

IMMUTEP LIMITED

By: /s/ Marc Voigt
Name: Marc Voigt
Title: Chief Executive Officer

Exhibit 99.1

Immutep Announces Initiation of TACTI-004 Phase III Trial in First Line Non-Small Cell Lung Cancer

SYDNEY, AUSTRALIA – December 10, 2024 – Immutep Limited (ASX: IMM; NASDAQ: IMMP) ("Immutep" or "the Company"), a clinical-stage biotechnology company developing novel LAG-3 immunotherapies for cancer and autoimmune disease, today announces the initiation of the pivotal TACTI-004 Phase III clinical trial for the treatment of first-line metastatic non-small cell lung cancer (1L NSCLC).

"The receipt of regulatory approval from the Australian Therapeutic Goods Administration to commence the TACTI-004 trial is a significant milestone for Immutep and marks its transformation into a Phase III company. This also represents a key step towards potentially establishing a new standard of care for patients with metastatic NSCLC. We are confident based on the strength of eftilagimod alfa's data that it can make a meaningful difference in cancer patients' lives, and we eagerly anticipate enrolling the first patient into this important study during the first quarter of 2025," said Marc Voigt, CEO of Immutep.

Immutep has successfully completed regulatory submissions to the vast majority of the more than 25 countries that will be part of the global TACTI-004 trial. Australia represents the first approval by all regulatory authorities including ethics committees and Institutional Review Boards (IRB). The Company also anticipates full approval in the United Kingdom shortly as it has received clearances from the Medicines and Healthcare products Regulatory Agency (MHRA) and the Research Ethics Committee (REC). Additional approvals from multiple countries are expected in the weeks and months ahead.

The registrational TACTI-004 Phase III trial will evaluate eftilagimod alfa, a soluble LAG-3 protein that activates dendritic cells, in combination with MSD's (Merck & Co., Inc., Rahway, NJ, USA) anti-PD-1 therapy KEYTRUDA® (pembrolizumab) and chemotherapy compared to KEYTRUDA in combination with chemotherapy and placebo in ~750 metastatic 1L NSCLC patients, regardless of PD-L1 expression. The 1:1 randomized, double-blind, multinational, controlled study, with dual primary endpoints of progression-free survival and overall survival, will include over 150 clinical sites in over 25 countries across the globe.

The Company expects to enrol the first patient in Q1 of CY2025.

KEYTRUDA® is a registered trademark of Merck Sharp & Dohme LLC, a subsidiary of Merck & Co., Inc., Rahway, NJ, USA.

About Immutep

Immutep is a clinical-stage biotechnology company developing novel LAG-3 immunotherapy for cancer and autoimmune disease. We are pioneers in the understanding and advancement of therapeutics related to Lymphocyte Activation Gene-3 (LAG-3), and our diversified product portfolio harnesses its unique ability to stimulate or suppress the immune response. Immutep is dedicated to leveraging its expertise to bring innovative treatment options to patients in need and to maximise value for shareholders. For more information, please visit www.immutep.com.

Australian Investors/Media:

Catherine Strong, Sodali & Co.
+61 (0)406 759 268; catherine.strong@sodali.com

U.S. Media:

Chris Basta, VP, Investor Relations and Corporate Communications
+1 (631) 318 4000; chris.basta@immutep.com

This announcement was authorised for release by the CEO of Immutep Limited.

Immutep Limited, Level 32, Australia Square, 264 George Street, Sydney NSW 2000, Australia
ABN: 90 009 237 889