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INNOCARE
诺诚健华

InnoCare Pharma Limited
諾誠健華醫藥有限公司
(Incorporated in the Cayman Islands with limited liability)
(Stock code: 9969)

INSIDE INFORMATION ANNOUNCEMENT

APPROVAL OF MINJUVI® (TAFASITAMAB) IN COMBINATION WITH LENALIDOMIDE FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY DIFFUSE LARGE B-CELL LYMPHOMA IN CHINA

This announcement is made by InnoCare Pharma Limited (the "**Company**") pursuant to Rule 13.09(2)(a) of the Rules Governing the Listing of Securities on The Stock Exchange of Hong Kong Limited (the "**Listing Rules**") and the inside information provision (as defined in the Listing Rules) under Part XIVA of the Securities and Futures Ordinance (Cap. 571 of the Laws of Hong Kong).

The board of directors of the Company (the "**Board**") is pleased to announce that the National Medical Products Administration (NMPA) has granted approval for Minjuvi® (tafasitamab), a humanized Fc-modified cytolytic CD19 targeting monoclonal antibody, in combination with lenalidomide, followed by Minjuvi monotherapy, for the treatment of adult patients with relapsed or refractory diffuse large B-cell lymphoma (DLBCL) who are not eligible for autologous stem cell transplant (ASCT). This is the first CD19 antibody approved for the treatment of relapsed or refractory DLBCL in China.

Tafasitamab, a humanized Fc-modified cytolytic CD19-targeting immunotherapy, in combination with lenalidomide, has already been approved for the treatment of eligible DLBCL patients in the region of Hong Kong, Macau and Taiwan. Furthermore, under the early access program in the Bo'ao Lecheng International Medical Tourism Pilot Zone and the Guangdong-Hong Kong-Macao Greater Bay Area, prescriptions of tafasitamab in combination with lenalidomide have been issued at Ruijin Hainan Hospital and Guangdong Clifford Hospital for eligible DLBCL patients.

Tafasitamab is approved under accelerated approval by the U.S. Food and Drug Administration (FDA), and conditionally approved by the European Medicines Agency (EMA), in combination with lenalidomide for the treatment of relapsed or refractory DLBCL adult patients who are not eligible for ASCT.

DLBCL is the most common type of non-Hodgkin lymphoma (NHL), and its incidence accounts for 31% to 34% of NHL globally. In China, DLBCL accounts for 45.8% of all NHL cases, according to the Hematology Branch of the Chinese Medical Association.

About Tafasitamab

Tafasitamab is a humanized Fc-modified cytolytic CD19 targeting monoclonal antibody. Tafasitamab incorporates an engineered Fc domain, which mediates B-cell lysis through apoptosis and immune effector mechanism including Antibody-Dependent Cell-Mediated Cytotoxicity (ADCC) and Antibody-Dependent Cellular Phagocytosis (ADCP).

MorphoSys and Incyte entered into: (a) in January 2020, a collaboration and licensing agreement to develop and commercialize tafasitamab globally; and (b) in February 2024, an agreement whereby Incyte obtained exclusive rights to develop and commercialize tafasitamab globally.

In August 2021, Incyte entered into a collaboration and license agreement with InnoCare for the development and exclusive commercialization of tafasitamab in hematology and oncology in Greater China.

In the United States, Monjuvi® (tafasitamab-cxix) received accelerated approval by the U.S. Food and Drug Administration in combination with lenalidomide for the treatment of adult patients with relapsed or refractory DLBCL not otherwise specified, including DLBCL arising from low grade lymphoma, and who are not eligible for ASCT. In Europe, Minjuvi® (tafasitamab) received conditional Marketing Authorization from the European Medicines Agency in combination with lenalidomide, followed by Minjuvi monotherapy, for the treatment of adult patients with relapsed or refractory DLBCL who are not eligible for ASCT.

Monjuvi, Minjuvi, the Minjuvi and Monjuvi logos and the “triangle” design are registered trademarks of Incyte.

Cautionary Statement as required by Rule 18A.08(3) of the Rules Governing the Listing of Securities on The Stock Exchange of Hong Kong Limited: There is no assurance that the Company will ultimately commercialize tafasitamab successfully. Shareholders and potential investors of the Company are advised to exercise caution when dealing in the shares of the Company.

By Order of the Board
InnoCare Pharma Limited
Dr. Jisong Cui
Chairperson and executive Director

Hong Kong, 21 May 2025

As at the date of this announcement, the Board of Directors comprises Dr. Jisong Cui as Chairperson and executive Director, Dr. Renbin Zhao as executive Director, Dr. Yigong Shi and Mr. Ronggang Xie as non-executive Directors, and Ms. Lan Hu, Dr. Dandan Dong and Prof. Kunliang Guan as independent non-executive Directors.