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Simcere Pharmaceutical Group Limited

先聲藥業集團有限公司

(Incorporated in Hong Kong with limited liability)

(Stock code: 2096)

VOLUNTARY ANNOUNCEMENT

THE CLINICAL TRIAL APPROVAL FOR SIM0500 FOR INJECTION (A HUMANIZED GPRC5D-BCMA-CD3 TRISPECIFIC ANTIBODY) ISSUED BY THE NATIONAL MEDICAL PRODUCTS ADMINISTRATION

This announcement is made by Simcere Pharmaceutical Group Limited (the “**Company**”, together with its subsidiaries, the “**Group**”) on a voluntary basis to inform shareholders and potential investors of the Company about the latest business development of the Group.

The board (the “**Board**”) of directors (the “**Directors**”) of the Company is pleased to announce that, on March 12, 2024, SIM0500 for injection, a humanized GPRC5D-BCMA-CD3 trispecific antibody, which is a new investigational anti-tumor drug independently developed by the Group, has obtained the Clinical Trial Approval issued by the National Medical Products Administration (國家藥品監督管理局) of China, which is intended to be investigated in a clinical trial for relapsed or refractory multiple myeloma.

ABOUT SIM0500

SIM0500 is a humanized GPRC5D-BCMA-CD3 trispecific antibody, which is a potential best-in-class (BIC) drug for the treatment of multiple myeloma based on the preclinical data. Through the research and development platform of multispecific antibody drugs with the Group’s own T-cell engagers, SIM0500 is a tumor-targeted T-cell activating drug, composed with the Group’s self-developed CD3 antibody with the feature activated by low affinity and high target activation and the antibody with anti-tumor associated antigen. It has the advantages of excellent tumor-killing effect and good tolerance. SIM0500 can potentially

overcome the drug resistance caused by the existing treatments, and show excellent anti-tumor activity in various animal pharmacodynamic models with different expression levels and has multiple advantages such as low effective dose and no recurrence of tumors after drug withdrawal.

On March 9, 2024, the Investigational New Drug (IND) application of SIM0500 has been approved by the U.S. Food and Drug Administration (FDA).

ABOUT THE COMPANY

The Company is an innovation and R&D-driven pharmaceutical company and has established a “State Key Laboratory of Neurology and Oncology Drug Development”. The Company focuses on the therapeutic areas of oncology, nervous system, autoimmune and anti-infection, with forward-looking layout of disease areas that may have significant clinical needs in the future, aiming to achieve the mission of “providing today’s patients with medicines of the future”. Driven by its in-house R&D efforts and synergistic innovation, the Company has established strategic cooperation partnerships with many innovative companies and research institutes.

By order of the Board
Sincere Pharmaceutical Group Limited
Mr. Ren Jinsheng
Chairman and Chief Executive Officer

Hong Kong, March 12, 2024

As at the date of this announcement, the Board comprises Mr. REN Jinsheng as the Chairman and executive Director, Mr. TANG Renhong, Mr. WAN Yushan and Ms. WANG Xi as the executive Directors; and Mr. SONG Ruilin, Mr. WANG Jianguo, Mr. WANG Xinhua and Mr. SUNG Ka Woon as the independent non-executive Directors.