

*Hong Kong Exchanges and Clearing Limited and The Stock Exchange of Hong Kong Limited take no responsibility for the contents of this announcement, make no representation as to its accuracy or completeness and expressly disclaim any liability whatsoever for any loss howsoever arising from or in reliance upon the whole or any part of the contents of this announcement.*



**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 SIM0613  
(LRRC15 ANTIBODY-DRUG CONJUGATE) 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 regarding the latest business development of the Group.

The board (the “**Board**”) of directors (the “**Directors**”) of the Company is pleased to announce that, SIM0613 for Injection, an LRRC15-targeting antibody-drug conjugate (“**ADC**”) independently developed by the Group, has received the Clinical Trial Approval issued by the National Medical Products Administration (國家藥品監督管理局) of China, and is intended for commencing the clinical trial for advanced solid tumors.

**ABOUT SIM0613**

SIM0613 targets the leucine-rich repeat-containing 15 (LRRC15), a protein highly expressed on various tumor types and cancer-associated fibroblasts but with limited expression on normal cells. Upon binding to the LRRC15 protein, SIM0613 is internalized where the cytotoxic payload is released, killing the cancer cell and therefore sparing healthy cells. SIM0613 is specifically engineered for deep tumor penetration, killing cancer cells and cancer-associated fibroblast cells, resulting in robust tumor regressions in multiple in vivo preclinical models.

In December 2025, a subsidiary of the Company, Jiangsu Simcere Zaiming Pharmaceutical Co., Ltd. (江蘇先聲再明醫藥有限公司) (“**Simcere Zaiming**”) entered into an exclusive licensing agreement (the “**License Agreement**”) with Ipsen Pharma SAS. (Euronext: IPN; ADR: IPSEY) (“**Ipsen**”). Under the terms of the License Agreement, Simcere Zaiming is eligible to receive up to US\$1.06 billion comprising upfront, development, regulatory and commercial milestone payments, and tiered royalties on sales, contingent upon successful development and regulatory approvals. Ipsen will have manufacturing rights, following the tech transfer process and will assume responsibility for all activities outside Greater China including Phase I preparation activities and submission of the Investigational New Drug and Clinical Trial applications.

## **ABOUT THE GROUP**

The Company is an innovation and R&D-driven pharmaceutical company and the Group has established a “State Key Laboratory of Neurology and Oncology Drug Development”. The Group focuses on the therapeutic areas of neuroscience, anti-oncology, 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 “for patients, for life”. 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  
**Simcere Pharmaceutical Group Limited**  
**Mr. Ren Jinsheng**  
*Chairman and Executive Director*

Hong Kong, May 6, 2026

*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; Mr. SONG Ruilin, Mr. WANG Jianguo, Mr. WANG Xinhua and Mr. SUNG Ka Woon as the independent non-executive Directors.*