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FOSUN PHARMA
復星医药

上海復星醫藥（集團）股份有限公司

Shanghai Fosun Pharmaceutical (Group) Co., Ltd.*

(a joint stock company incorporated in the People's Republic of China with limited liability)

(Stock Code: 02196)

OVERSEAS REGULATORY ANNOUNCEMENT

This announcement is made pursuant to Rule 13.10B of the Rules Governing the Listing of Securities on The Stock Exchange of Hong Kong Limited.

The following sets out the "Announcement in Relation to the Progress of Drug Clinical Trial By a Subsidiary" published by Shanghai Fosun Pharmaceutical (Group) Co., Ltd.* (the "**Company**") on the website of the Shanghai Stock Exchange, for your reference only. The following is a translation of the above mentioned announcement solely for the purpose of providing information. Should there be any discrepancies, the Chinese version will prevail.

By order of the Board
Shanghai Fosun Pharmaceutical (Group) Co., Ltd.*
Chen Yuqing
Chairman

Shanghai, the PRC
4 November 2026

As at the date of this announcement, the executive directors of the Company are Mr. Chen Yuqing, Ms. Guan Xiaohui, Mr. Wen Deyong, Mr. Wang Kexin and Mr. Liu Yi; the non-executive directors of the Company are Mr. Chen Qiyu and Mr. Pan Donghui; the independent non-executive directors of the Company are Mr. Yu Tze Shan Hailson, Mr. Wang Quandi, Mr. Chen Penghui and Mr. Yang Yucheng; and the employee director of the Company is Ms. Yan Jia.

* *for identification purposes only*

Shanghai Fosun Pharmaceutical (Group) Co., Ltd.* Announcement in Relation to the Progress of Drug Clinical Trial By a Subsidiary

The board of directors of the Company and all directors warrant that this announcement does not contain any false information, misleading statement or material omission, and accept legal liability for the truthfulness, accuracy and completeness of the contents herein contained.

I. Overview

Recently, Shanghai Fosun Pharmaceutical Industrial Development Company Limited* (上海復星醫藥產業發展有限公司), a subsidiary of Shanghai Fosun Pharmaceutical (Group) Co., Ltd.*（上海復星醫藥（集團）股份有限公司）(the "**Company**"), initiated the Phase III clinical trial of FXS6837 capsules (former development code: XH-S003, "**FXS6837**") for the treatment of paroxysmal nocturnal hemoglobinuria ("**PNH**") in Chinese mainland[1].

II. Basic Information and Research Progress of FXS6837

FXS6837 is a small molecule inhibitor of complement system with independent intellectual property rights owned by the Group (i.e., the Company and its subsidiaries/units, the same applies below), and is intended to be used for the treatment of diseases with abnormal complement system activation.

As of the date of this announcement (i.e., 4 November 2026), another indication of FXS6837 (for the treatment of glomerular diseases with abnormal complement activation such as IgA nephropathy) is at the stage of Phase IIb clinical trial in Chinese mainland.

[1] Excluding Hong Kong, Macau and Taiwan regions for the purpose of this announcement, the same applies below.

As of September 2026, the Group had invested approximately RMB199 million (unaudited) in total in the research and development of FXS6837 at this stage.

According to the latest data from IQVIA CHPA[2], in 2025, the sales of key drugs for the treatment of PNH in Chinese mainland was approximately RMB295 million.

III. Risk Reminder

As required by the relevant regulations, FXS6837 is still required to undergo relevant clinical trials in Chinese mainland and obtain approval from the national drug review authority before it can be launched.

The process from drug development and clinical research to launch involves a long period and multiple stages, with various uncertainties. Investors should take note of the investment risks.

Announcement is hereby made.

Board of Directors of Shanghai Fosun Pharmaceutical (Group) Co., Ltd.*

4 November 2026

* For identification purpose only

[2] Provided by IQVIA, a global provider of professional medical and health information and strategic consultation service; IQVIA CHPA data covers the drug sales market of hospitals with more than 100 beds in Chinese mainland, the actual sales of different drugs may vary from the IQVIA CHPA data to varying degrees due to their different sales channels.