FOSUN PHARMA
复星医药

上海復星醫藥（集團）股份有限公司

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

(a joint stock limited 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 Approval for Drug Registration of 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 abovementioned 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.*
Wu Yifang
Chairman

Shanghai, the PRC
24 April 2024

As at the date of this announcement, the executive directors of the Company are Mr. Wu Yifang, Mr. Wang Kexin, Ms. Guan Xiaohui and Mr. Wen Deyong; the non-executive directors of the Company are Mr. Chen Qiyu, Mr. Yao Fang, Mr. Xu Xiaoliang and Mr. Pan Donghui; and the independent non-executive directors of the Company are Ms. Li Ling, Mr. Tang Guliang, Mr. Wang Quandi and Mr. Yu Tze Shan Hailson.

* *for identification purposes only*

Shanghai Fosun Pharmaceutical (Group) Co., Ltd.*
Announcement in Relation to the Approval for Drug Registration of 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

Shenyang Hongqi Pharmaceutical Co., Ltd.（瀋陽紅旗製藥有限公司）, 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**" and, together with its subsidiaries/units, the "**Group**"), recently received the marketing registration approval from the National Medical Products Administration for the independently researched and developed Cycloserine Capsule (the "**New Drug**") for the treatment of active tuberculosis and extrapulmonary tuberculosis (including renal tuberculosis) caused by tuberculosis bacteria that is susceptible to the drug and that are not effectively treated with first-line anti-tuberculosis drugs (such as streptomycin, isoniazid, rifampin, and ethambutol).

II. General Information of the New Drug

Generic Name: Cycloserine capsule

Dosage Form: Capsules

Specification: 0.25g

Registration Category: Chemicals Category 3

Marketing Authorization Holder/ Pharmaceutical Manufacturer: Shenyang Hongqi Pharmaceutical Co., Ltd.

Drug Approval Number: Guo Yao Zhun Zi H20243531

III. Research and Marketing Progress of the New Drug

The New Drug is a chemical drug independently developed by the Group, which is mainly used for the treatment of active tuberculosis and extrapulmonary tuberculosis (including renal tuberculosis) caused by tuberculosis bacteria that is susceptible to the drug and that are not effectively treated with first-line anti-tuberculosis drugs (such as streptomycin, isoniazid, rifampin, and ethambutol).

As of March 2024, the Group has invested approximately RMB11.76 million (unaudited) in total in the research and development of the New Drug at the current stage.

As at the date of this announcement, in addition to the New Drug, other cycloserine capsules which have been approved to be launched in China (excluding Hong Kong, Macau, and Taiwan Region for the purpose of this announcement, the same applies below) include cycloserine capsules of Dong-A ST Co., Ltd. and Zhejiang Hisun Pharmaceutical Co., Ltd. * (浙江海正藥業股份有限公司). According to the latest data from IQVIA CHPA[1], the sales of the cycloserine capsule in China in 2023 was approximately RMB 300 million.

IV. Impact on the Listed Company and Risk Warning

The approval for marketing registration of the New Drug will bring more options for the medication for active tuberculosis and extrapulmonary tuberculosis (including renal tuberculosis) and will further enrich the product line of the Group. It is expected that the approval of the New Drug will not have a material impact on the Group's results at this stage.

Due to the characteristics of the pharmaceutical products industry, the specific sales performance of pharmaceutical products after launch may be affected by factors including, but not limited to, the demand for medication, market competition and sales channels, and is subject to considerable uncertainty. Investors should take note of the investment risks.

[1] provided by IQVIA, a provider of professional medical and health information and strategic consultation service in the world; IQVIA CHPA data represents the drug sales market of hospitals with more than 100 beds in China, the actual sales of different drugs may vary from the IQVIA CHPA data to varying degrees due to their different sales channels.

Announcement is hereby made.

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

24 April 2024

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