上海復旦張江生物醫藥股份有限公司

Shanghai Fudan-Zhangjiang Bio-Pharmaceutical Co., Ltd.*

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

(Stock code:1349)

INDICATIVE ANNOUNCEMENT
FIRST PATIENT ENROLLED IN CONFIRMATORY CLINICAL TRIAL OF AMINOLEVULINIC ACID HYDROCHLORIDE POWDER FOR ORAL SOLUTION FOR INTRAOPERATIVE VISUALISATION OF GLIOMA

This announcement is made by Shanghai Fudan-Zhangjiang Bio-Pharmaceutical Co., Ltd.* (the "**Company**", together with its subsidiaries, the "**Group**") on a voluntary basis.

The first patient in the confirmatory clinical trial of aminolevulinic acid hydrochloride powder for oral solution (the "**Drug**") for intraoperative visualisation of glioma (WHO grade III or IV) (the "**Study**") has recently been successfully enrolled in the Study.

ABOUT THE DRUG

Glioma is the most common primary intracranial tumor, which is characterized by high incidence, high recurrence rate, high mortality rate and low cure rate. Surgical resection is the stand of care at domestic and abroad, and the survival and prognosis of patients is related to the degree of surgical resection. Therefore, the basic principle of surgery is to remove as much diseased tissue as possible without damaging adjacent normal brain tissue. However, most of the gliomas are invasive growth. The boundary between gliomas and the surrounding normal brain tissue is not clear so that it is difficult to conduct complete surgical resection. The Company intends to develop this intraoperative fluorescence-guided technology to visualize the tumour margin, so as to guide the resection range in real time, to help surgeons improve complete resection rate while reserving healthy tissue. The Company wish this technology could improve the postoperative quality of life of patients and prolong the survival period of patients.

RESEARCH AND DEVELOPMENT INFORMATION AND PROGRESS OF THE DRUG

Taizhou Fudan-Zhangjiang Pharmaceutical Co., Ltd* (泰州復旦張江藥業有限公司), a subsidiary of the Company, has received the acceptance notice for the confirmatory clinical trial application in December 2023 which was approved in March 2024. The Study aims to evaluate its effectiveness and safety of aminolevulinic acid hydrochloride powder for oral solution (5-ALA) for fluorescence-guided tumour resection in patients with malignant high-grade glioma (WHO grade III or IV). As at the date on the publication of this announcement, the first patient has been successfully enrolled in the Study. After the completion of the clinical trial, the listing application of the Drug can be submitted in accordance with relevant laws and regulations.

Due to the high-tech, high-risk and high-value-added characteristics of pharmaceutical products, and the long cycle and numerous stages in the process, there are uncertainties in drug pre-clinical research, clinical trial and commercialization. These many stages make it susceptible to uncertainties and therefore, investors are advised to make cautious decisions and pay careful attention to investment risks. The Company will actively facilitate the above research and development project and fulfill its information disclosure obligations in a timely manner for subsequent progress in strict accordance with relevant regulations.

By order of the Board
Zhao Da Jun
Chairman

As at the date on the publication of this announcement, the Board comprises:

Mr. Zhao Da Jun (Executive Director)
Ms. Xue Yan (Executive Director)
Mr. Shen Bo (Non-executive Director)
Ms. Yu Xiao Yang (Non-executive Director)
Mr. Wang Hong Guang (Independent Non-executive Director)
Mr. Lam Siu Wing (Independent Non-executive Director)
Mr. Xu Pei Long (Independent Non-executive Director)

Shanghai, the PRC

20 June 2024
** For identification purpose only*