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(Incorporated in the Cayman Islands with limited liability)

(Stock Code: 2005)

VOLUNTARY ANNOUNCEMENT UPDATE ON PRODUCT DEVELOPMENT

The board of directors (the "Board") of SSY Group Limited (the "Company", together with its subsidiaries, the "Group") is pleased to announce that the Group has obtained the approval for drug production and registration for Allopurinol Tablets (0.1g) from the National Medical Products Administration of China (the "NMPA"), being under type 3 chemical drug and regarded as passing the consistency evaluation.

Allopurinol Tablet is mainly used for patients with primary or secondary gout, for patients with leukemia, lymphoma and malignant tumors undergoing treatment, and for the treatment of patients with recurrent calcium oxalate stones. As stated in the Company's announcement dated 28 April 2025, the Group's Allopurinol bulk drug has obtained the approval for registration from the NMPA to become a bulk drug for the preparations on the market.

This announcement is a voluntary announcement made by the Company to keep the shareholders and potential investors informed of the latest business development of the Group.

By order of the Board
Chow Hing Yeung
Executive Director and Company Secretary

Hong Kong, 1 September 2025

As at the date of this announcement, the Board comprises Mr. Qu Jiguang, Mr. Su Xuejun, Mr. Meng Guo, Mr. Chow Hing Yeung and Ms. Qu Wanrong as executive Directors, Mr. Liu Wenjun as non-executive Director, and Mr. Wang Yibing, Mr. Chow Kwok Wai and Mr. Jiang Guangce as independent non-executive Directors.