Hansoh Pharmaceutical Group Company Limited
翰森製藥集團有限公司

(Incorporated in the Cayman Islands with limited liability)

(Stock Code: 3692)

VOLUNTARY ANNOUNCEMENT

AUMOLERTINIB MESYLATE TABLETS WAS APPROVED IN THE EU AS MONOTHERAPY

The board of directors (the "**Board**") of Hansoh Pharmaceutical Group Company Limited (the "**Company**" and together with its subsidiaries, the "**Group**") is pleased to announce that, on February 12, 2026, Aumolertinib Mesylate Tablets, marketed as Ameile (阿美樂®) in China and Aumseqa® outside China, the Group's innovative drug, has been approved in the European Union (the "**EU**") as monotherapy for: (i) the first-line treatment of adult patients with advanced non-small cell lung cancer ("**NSCLC**") whose tumours have epidermal growth factor receptor ("**EGFR**") exon 19 deletions or exon 21 (L858R) substitution mutations; and (ii) the treatment of adult patients with advanced EGFR T790M mutation-positive NSCLC. The approval by the European Commission (EC) follows the positive opinion of the Committee for Medicinal Products for Human Use (CHMP) of the European Medicines Agency (EMA).

About Aumolertinib Mesylate Tablets

Aumolertinib Mesylate Tablets is the first original third-generation EGFR-TKI innovative drug in China. It has been approved for five indications by the National Medical Products Administration (NMPA), namely: it was approved for the treatment of patients with locally advanced or metastatic NSCLC with T790M mutation, who have progressed on or after EGFR-TKI therapy in 2020; it was approved as the first-line treatment for adult patients with locally advanced or metastatic NSCLC whose tumors have EGFR exon 19 deletions or exon 21 (L858R) substitute mutation positive in 2021; it was approved for the treatment of patients with locally advanced, unresectable NSCLC whose disease has not progressed following definitive platinum-based chemoradiotherapy whose tumors have EGFR exon 19 deletions or exon 21 (L858R) substitute mutations in 2025; it was approved for the adjuvant treatment of adult patients with stage II to IIIB NSCLC whose tumors have EGFR exon 19 deletions or exon 21 L858R mutations, and who have undergone tumor resection with or without prior adjuvant chemotherapy as determined by their physician in 2025; it was approved that Ameile in combination with pemetrexed and platinum-based chemotherapy to be used as the first-line treatment of adult patients with locally advanced or metastatic NSCLC whose tumors have EGFR exon 19 deletions or exon 21 (L858R) mutations in 2026. Additionally, Aumolertinib Mesylate Tablets was approved by the Medicines and Healthcare Products Regulatory Agency in the United Kingdom (MHRA) for marketing in 2025.

By Order of the Board
Hansoh Pharmaceutical Group Company Limited
Zhong Huijuan
Chairlady

Hong Kong, February 20, 2026

As at the date of this announcement, the Board comprises Ms. Zhong Huijuan as chairlady and executive director, Ms. Sun Yuan and Dr. Lyu Aifeng as executive directors, and Mr. Lin Guoqiang, Mr. Chan Charles Sheung Wai, Ms. Yang Dongtao and Mr. Yan Jia as independent non-executive directors.