December 16, 2025

To,
Dy. General Manager
Department of Corporate Services,
BSE Ltd.
Phiroze Jeejeebhoy Towers
Dalal Street, Fort, Mumbai – 400 001

Ref: Scrip Code: 532296

To,
The Manager – Listing,
The National Stock Exchange of India Ltd.,
Plot No. C/1, G Block
Bandra Kurla Complex,
Bandra (E), Mumbai – 400 051

Ref: Scrip Name: GLENMARK

Dear Sir,

Sub: Glenmark Secures Exclusive Multi-Regional Rights to Aumolertinib, an MHRA and NMPA approved Third-Generation EGFR-TKI, from Hansoh Pharma

Glenmark Specialty S.A. (GSSA), a wholly owned subsidiary of Glenmark Pharmaceuticals Ltd. (Glenmark), today announced that it has entered into an exclusive license, collaboration and distribution agreement with Jiangsu Hansoh Pharmaceutical Group Co., Ltd. (Hansoh Pharma) for Aumolertinib, a third-generation Epidermal Growth Factor Receptor Tyrosine Kinase Inhibitor (EGFR-TKI) for the treatment of non-small cell lung cancer (NSCLC).

Kindly find attached media release which is self- explanatory.

Request you to kindly take the same on record.

Thanking you,

Yours faithfully,
For Glenmark Pharmaceuticals Limited

HARISH VINAYAK KUBER
Digitally signed by HARISH VINAYAK KUBER
Date: 2025.12.16 15:33:50 +05'30'

Harish Kuber
Company Secretary & Compliance Officer

Encl: As above

Press Release
For Immediate Distribution

Glenmark Secures Exclusive Multi-Regional Rights to Aumolertinib, an MHRA and NMPA approved Third-Generation EGFR-TKI, from Hansoh Pharma

Partnership strengthens Glenmark's oncology strategy across high-potential markets

Neuchatel, Switzerland and Mumbai, India, December 16, 2025: Glenmark Specialty S.A. (GSSA), a wholly owned subsidiary of Glenmark Pharmaceuticals Ltd. (Glenmark), today announced that it has entered into an exclusive license, collaboration and distribution agreement with Jiangsu Hansoh Pharmaceutical Group Co., Ltd. (Hansoh Pharma) for Aumolertinib, a third-generation Epidermal Growth Factor Receptor Tyrosine Kinase Inhibitor (EGFR-TKI) for the treatment of non-small cell lung cancer (NSCLC).

Under the terms of the agreement, Glenmark receives exclusive rights to develop and commercialize Aumolertinib across its licensed territories: Middle East and Africa, Southeast & South Asia, Australia, New Zealand, Russia/CIS and a few selected Caribbean countries covered by the agreement. Hansoh Pharma will receive an upfront payment of low double-digit million USD, followed by potential regulatory and commercial milestone payments possibly cumulating to over US$1 billion, in addition to tiered royalties on net sales in the licensed territories.

"At Glenmark, we remain focused on building a growth-oriented oncology business across high-potential markets. Aumolertinib is a strategic addition that strengthens our near-term commercial pipeline and enhances our ability to serve patients with EGFR-mutated lung cancer. This collaboration reinforces our disciplined approach to expanding our innovative portfolio and widening access to advanced cancer care across key markets," **said Glenn Saldanha, Chairman and Managing Director, Glenmark Pharmaceuticals Ltd.**

Aumolertinib, (marketed as Ameile® in China and Aumseqa® in the United Kingdom and Europe), as a monotherapy, has received marketing authorization from the UK MHRA and is indicated the first-line treatment of adult patients with locally advanced or metastatic non-small cell lung cancer ("NSCLC") with activating epidermal growth factor receptor ("EGFR") mutations, and the treatment of adult patients with locally advanced or metastatic EGFR T790M mutation-positive NSCLC. It has also received approval for four indications in China (second-line T790M mutation, first-line NSCLC EGFR mutated, unresectable Stage III post-chemoradiotherapy, and adjuvant Stage II–IIIB NSCLC). Aumolertinib became Hansoh Pharma's first innovative drug approved in an overseas market and the first China-developed EGFR-TKI to be launched internationally.

About Aumolertinib

Ameile (Aumolertinib Mesilate Tablets) is the first original third-generation EGFR-TKI innovative drug in China. It has been approved for four indications in China, namely: in March 2020, 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 December 2021, 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 March 2025, 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 May 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. Additionally, in June 2025, Aumolertinib (trade name in the United Kingdom:

Aumseqa®) was approved by the Medicines and Healthcare Products Regulatory Agency in the United Kingdom (MHRA) for marketing.

About Glenmark Pharmaceuticals Limited

Glenmark Pharmaceuticals Ltd. (BSE: 532296 | NSE: GLENMARK) is a research-led, global pharmaceutical company, having a presence across Branded, Generics, and OTC segments; with a focus on therapeutic areas of respiratory, dermatology and oncology. The company has 11 world-class manufacturing facilities spread across 4 continents, and operations in over 80 countries. Scrip 100 positions Glenmark amongst the Top 100 biopharmaceutical companies ranked by Pharmaceutical Sales in 2023. Glenmark's Green House Gas (GHG) emission reduction targets have been approved in 2023 by the Science Based Target initiative (SBTi), making it only the second pharmaceutical company in India to achieve this. The organization has impacted over 4.5 million lives over the last decade through its CSR interventions. For more information, visit www.glenmarkpharma.com. You can follow us on LinkedIn (Glenmark Pharmaceuticals) and Instagram (Glenmark_pharma).

About Hansoh Pharma

Hansoh Pharma is a leading innovation-driven pharmaceutical enterprise headquartered in China. With the mission of "continuous innovation for better life", the company focuses on major disease therapeutic areas such as oncology, anti-infectives, central nervous system (CNS), metabolism and autoimmunity. Hansoh Pharma has launched 7 innovative drugs that generate product sales in China, with the revenue from innovative drugs and collaborative products exceeding 80%, forming a rich product pipeline. The company has consistently ranked among the top 100 global pharmaceutical companies and is recognized as one of the top 3 pharmaceutical R&D enterprises in China, and is designated as a National Key High-Tech Enterprise and a National Technology Innovation Demonstration Enterprise. Hansoh Pharma was listed on the Hong Kong Stock Exchange in June 2019 (stock code: 03692.HK).

For more information, please visit www.hspharm.com.

For more information, please contact
Aakanksha Pilay | mediarelations@glenmarkpharma.com | +91 7218171062
Adfactors PR | glenmark@adfactorspr.com | +91 88502 91322