June 23, 2021

To,
Dy. General Manager
Department of Corporate Services,
BSE Ltd.,
P. J. 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 Pharmaceuticals receives ANDA approval for Arformoterol Tartrate Inhalation Solution, 15 mcg/2 mL, Unit-Dose Vials

With reference to the subject mentioned above, 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: 2021.06.23 08:31:31 +05'30'

Harish Kuber
Company Secretary & Compliance Officer

Encl: as above

Press Release **For Immediate Release**

Glenmark Pharmaceuticals receives ANDA approval for Arformoterol Tartrate Inhalation Solution, 15 mcg/2 mL, Unit-Dose Vials

Mumbai, India; June 23, 2021: Glenmark Pharmaceuticals Ltd (Glenmark) has received final approval by the United States Food & Drug Administration (U.S. FDA) for Arformoterol Tartrate Inhalation Solution, 15 mcg/2 mL, Unit-Dose Vials, the generic version of BROVANA®[1] Inhalation Solution, 15 mcg/2 mL, of Sunovion Pharmaceuticals Inc. Arformoterol Tartrate Inhalation Solution, 15 mcg/2 mL, Unit-Dose Vials will be manufactured in the company's North American manufacturing facility based in Monroe, North Carolina, and marks the company's first nebulizer approval.

According to IQVIA™ sales data for the 12 month period ending April 2021, the BROVANA® Inhalation Solution, 15 mcg/2 mL market[2] achieved annual sales of approximately $437.9 million*.

Commenting on the launch, Sanjeev Krishan, President, Glenmark North America said, "We are very excited to be one of the first generic companies to receive approval for such an important product for our customers. This also marks our third approval from our state-of-the art manufacturing facility in Monroe in 2021, demonstrating our capability to offer high quality medicines with affordable access across multiple dosage platforms."

Glenmark's current portfolio consists of 172 products authorized for distribution in the U.S. marketplace and 44 ANDA's pending approval with the U.S. FDA. In addition to these internal filings, Glenmark continues to identify and explore external development partnerships to supplement and accelerate the growth of its existing pipeline and portfolio.

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About Glenmark Pharmaceuticals Ltd

Glenmark Pharmaceuticals Ltd. (GPL) is a global research-led pharmaceutical company with presence across Generics, Specialty and OTC business with operations in over 50 countries. Glenmark's key therapy focus areas globally are respiratory, dermatology and oncology. It ranks among the world's top 50 Generics and Biosimilars companies (Top 50 Company Rankings, 2020, from Informa's Generics Bulletin). The company has been listed in the Dow Jones Sustainability Index (DJSI), under the category of emerging markets for the third consecutive year in a row. DJSI is one of the world's most respected and widely accepted sustainability benchmarks globally with only the top ranked companies in terms of Corporate Sustainability within each industry are featured in the index. For more information, visit www.glenmarkpharma.com

For more information, please contact:

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[1]All brand names and trademarks are the property of their respective owners.

[2]Market includes brand and all available therapeutic equivalents

*IQVIA™ National Sales Perspectives: Retail & Non-Retail, April 2021