June 21, 2023

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,

Re: Disclosure under Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015

Subject: Update on the Company's Monroe, North Carolina (USA) Facility

We refer to our letter dated 23rd August 2022 informing you about the "Official Action Indicated" OAI status of the Company's Monroe, North Carolina (USA) manufacturing facility by US FDA following the US FDA inspection conducted between April 04 to May 19, 2022. We wish to inform you that the US FDA has now issued a warning letter to the Monroe facility.

The Company had done a voluntary recall of all its products from this site in August 2021 and since then has not been commercializing any product from this site. Hence, the warning letter will have no impact on the existing revenues.

Glenmark continues to cooperate with the US FDA and is committed to undertake all necessary steps required to address their observations at the earliest. The Company is committed to maintaining the highest quality and compliant manufacturing standards at all of its facilities across the globe.

This may be considered as a disclosure pursuant to Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015.

This is for your information.

Thanking you,

Yours faithfully,
For Glenmark Pharmaceuticals Limited

HARISH VINAYAK KUBER
Digitally signed by HARISH VINAYAK KUBER
Date: 2023.06.21 20:41:54 +05'30'

Harish Kuber
Company Secretary & Compliance Officer

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