

September 21, 2024

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

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 Code: 532296 Ref: Scrip Name: GLENMARK

Dear Sir,

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

Sub.: U.S. FDA inspection at the Company's Facility at Chhatrapati Sambhaji Nagar (Aurangabad), India

This is to inform you that the United States Food & Drug Administration (USFDA) completed a Good Manufacturing Practice (GMP) inspection at the Company's formulation manufacturing facility based out of Chhatrapati Sambhaji Nagar (Aurangabad), India. The inspection was conducted from 9th September to 20th September, 2024. The inspection closed with no observation.

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 Kuber
Company Secretary & Compliance Officer