

August 21, 2026

To Listing Department, NATIONAL STOCK EXCHANGE OF INDIA LIMITED Exchange Plaza, Bandra Kurla Complex, Bandra (E), MUMBAI -400 051 Company Code No. AUOPHARMA	To The Corporate Relations Department BSE LIMITED Phiroz Jeejeebhoy Towers, 25 th floor, Dalal Street, MUMBAI -400 001 Company Code No. 524804
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Dear Sir / Madam,

Sub: Completion of US FDA Inspection at Auro Peptides Ltd, a subsidiary/business arm of the Company.

Pursuant to Regulation 30 of the SEBI (Listing Obligations & Disclosure Requirements) Regulations, 2015, this is to inform you that:

The United States Food and Drug Administration (US FDA) conducted a routine inspection at AuroPeptides Ltd, a manufacturer of Peptide APIs, located at Indrakaran Village, Kandi Mandal, Sangareddy District, Telangana - 502329, from 17th to 21st August, 2026.

We are pleased to inform you that the inspection concluded with only one (1) observation, relating to facility and equipment maintenance. This observation does not pertain to data integrity or related GMP compliance. Corrective action was already initiated during the course of the inspection, and our formal response will be submitted within the stipulated timeline.

This outcome reaffirms the Company's continued commitment to maintaining robust quality and compliance standards across its manufacturing facilities.

The Company will keep the Stock Exchanges informed of any further developments in this regard, as required under the applicable regulations.

Please take the above information on record.

Yours faithfully,

For AUROBINDO PHARMA LIMITED

B. Adi Reddy
Company Secretary

AUROBINDO PHARMA LIMITED

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(CIN : L24239TG1986PLC015190)

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