

August 28, 2026

To Listing Department, NATIONAL STOCK EXCHANGE OF INDIA LIMITED Exchange Plaza, Bandra Kurla Complex, Bandra (E), MUMBAI -400 051 Company Code No. AUROPHARMA	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 Unit-VI of Apitoria Pharma Private Limited, a wholly owned subsidiary of the Company – Reg.,

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

The United States Food and Drug Administration (US FDA) inspected Unit-VI, an API manufacturing facility of Apitoria Pharma Private Limited, a wholly owned subsidiary of the Company, situated at E-Bonangi Village, Parawada Mandal, Anakapalli District, 531019, Andhra Pradesh from August 24 to August 28, 2026.

We wish to inform you that the inspection concluded with three observations, which are procedural in nature, and we will respond to the US FDA within the stipulated timeline.

The Company is committed 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

(CIN : L24239TG1986PLC015190)

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