

May 6, 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: Classification of Unit-VII of Aurobindo Pharma Limited, Jedcherla as Voluntary Action Indicated (VAI) – Reg.,

Ref: Our letter dated February 10, 2026

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) inspected Aurobindo Pharma's Unit-VII, an oral solid dosage manufacturing unit, situated at Special Economic Zone (Pharma), TSIIIC, Green Industrial Park, Polepally Village, Jedcherla Mandal, Mahaboobnagar District, 509302, Telangana from January 28 to February 10, 2026. At the end of the inspection, a 'Form 483' was issued with a total of 9 observations.

The Unit has now received Establishment Inspection Report (EIR) classifying the facility as "Voluntary Action Indicated" ("VAI"). FDA has concluded that this inspection is now closed.

Please take the above information on record.

Yours faithfully,

For AUROBINDO PHARMA LIMITED

B. Adi Reddy
Company Secretary

AUROBINDO PHARMA LIMITED

www.aurobindo.com

(CIN : L24239TG1986PLC015190)

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