

September 30, 2023

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 Phiroze 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 our Unit VI - B – Reg.,

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 Unit VI - B, a Formulation manufacturing facility of the Company, situated at Chitkul Village, Patancheru Mandal, Sangareddy District, Telangana, from September 22 to September 29, 2023.

At the end of the inspection, a 'Form 483' was issued with 1 observation which is procedural in nature. We will respond to the US FDA within the stipulated timelines and work closely with US FDA to close the observation at the earliest.

We will keep the stock exchanges further informed if there is any material information relating to the above in the future.

Please take the above information on record.

Thanking you,

Yours faithfully,
For **AUROBINDO PHARMA LIMITED**

B. Adi Reddy
Company Secretary

AUROBINDO PHARMA LIMITED

(CIN : L24239TG1986PLC015190)

www.aurobindo.com

PAN No. AABCA7366H

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