

December 17, 2024

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

**Sub: Completion of US FDA Inspection at Unit V of Apitoria Pharma Private Limited, a wholly owned subsidiary of the Company – 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-V, an API manufacturing facility, of Apitoria Pharma Private Limited, a wholly owned subsidiary of the Company, situated at Pashamylaram Village, Patancheru Mandal, Sanga Reddy District, Telangana from December 09 to December 17, 2024.

The inspection closed with 02 observations. The observations are of procedural in nature and will be responded to within the stipulated time.

We will keep the stock exchanges informed if there is any further 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)

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