

December 20, 2023

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: Classification of US FDA Inspection at Unit IV of APL Healthcare Ltd. – Reg.,**

**Ref: Our letter dated September 20, 2023**

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

The U.S. Food and Drug Administration (FDA) had conducted an inspection at the Unit IV, a Formulation manufacturing facility, of APL Healthcare Limited, a wholly owned subsidiary of the Company, situated at Menakuru Village, Naidupeta Mandal, Tirupati District, Andhra Pradesh, from September 13 to September 19, 2023.

The Unit has now received Establishment Inspection Report classifying the facility as "Voluntary Action Indicated" ("VAI").

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](http://www.aurobindo.com)

PAN No. AABCA7366H

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