

March 11, 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> Phiroz 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: Acceptance of compliance with WHO GMP by WHO Prequalification Unit (PQT) Inspection Services Team (INS) for APL Healthcare Limited, Unit IV – Reg.,**

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

The Prequalification Unit Inspection Team (PQT-INS) of World Health organization (WHO) 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 11 to September 15, 2023.

Based on the inspection findings and our response to the same, the PQT-INS has decided to recommend to the Prequalification Assessment Team that the above facility be named as a manufacturing site in the dossier for Dolutegravir, Lamivudine, Tenofovir Disoproxil Fumarate Tablets (50/300/300 mg).

Please take the above information on record.

Thanking you,

Yours faithfully,  
**For AUROBINDO PHARMA LIMITED**

**B.Adi Reddy**  
**Company Secretary**

**AUROBINDO PHARMA LIMITED**  
[www.aurobindo.com](http://www.aurobindo.com)

(CIN : L24239TG1986PLC015190)

PAN No. AABCA7366H

Corp. Off.: Galaxy, Floors: 22-24, Plot No.1, Survey No.83/1, Hyderabad Knowledge City, Raidurg Panmaktha, Ranga Reddy District, Hyderabad – 500 032, Telangana, India.

Tel : +91 40 6672 5000 / 6672 1200 Fax: +91 40 6707 4044.

Regd.off.: Plot No.2, Maithrivihar, Ameerpet, Hyderabad-500038 T.S., INDIA Tel: +91 40 2373 6370 / 2374 7340 Fax: +91 40 2374 1080 / 2374 6833  
Email: [info@aurobindo.com](mailto:info@aurobindo.com) Website: [www.aurobindo.com](http://www.aurobindo.com)