

February 16, 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. AUOPHARMA</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 synthetic peptide API facility of Auro Peptides Ltd., a 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 manufacturing unit of Auro Peptides Ltd, the synthetic peptides API business arm of the company, situated at Indrakaran Village, Kandi Mandal, Sangareddy District, Telangana, was audited by the US FDA from 12<sup>th</sup> to 16<sup>th</sup> February 2024. The inspection was concluded with zero observations.

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)