

August 05, 2024

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: Classification of US FDA Inspection at Unit II of Eugia Pharma Specialities Ltd. (our wholly owned subsidiary) – Reg.,

Ref: Our letter dated May 03, 2024

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) had conducted an inspection at Unit-II, a Formulation manufacturing facility, of Eugia Pharma Specialities Ltd., a wholly owned subsidiary of the Company, situated at Bhiwadi, Alwar, Rajasthan, from April 25th to May 3rd 2024.

Subsequently, the US FDA has determined the inspection classification status of this facility as 'Official Action Indicated (OAI)'.

The company remains committed to work closely with the US FDA and continues to enhance its compliance on an ongoing basis.

We will keep the stock exchanges informed about further developments.

Please take the above information on record.

Thanking you,

Yours faithfully,
For **AUROBINDO PHARMA LIMITED**

B. Adi Reddy
Company Secretary

AUROBINDO PHARMA LIMITED

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(CIN : L24239TG1986PLC015190)

PAN No. AABCA7366H

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