November 29, 2023

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** Phiroz Jeejeebhoy Towers, 25th floor, Dalal Street, **MUMBAI -400 001** **Company Code No. 524804**

Dear Sir/ Madam,

Sub: Press Release - Aurobindo Pharma receives USFDA Approval for Darunavir Tablets, 600 mg and 800 mg.

We enclose a copy of the Press Release that is being issued by the Company in connection with USFDA approval received by the Company, for Darunavir Tablets, 600 mg and 800 mg.

Please take the information on record.

Thanking you,

Yours faithfully,
For **AUROBINDO PHARMA LIMITED**

BADDIGAM ADI REDDY
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Date: 2023.11.29 11:33:57 +05'30'

B. Adi Reddy
Company Secretary

Encl: as above

Hyderabad, India, November 29, 2023

Aurobindo Pharma receives USFDA Approval for Darunavir Tablets, 600 mg and 800 mg

Aurobindo Pharma Limited is pleased to announce that it has received final approval from the US Food & Drug Administration (USFDA) to manufacture and market Darunavir Tablets, 600 mg and 800 mg, which is bioequivalent and therapeutically equivalent to the reference listed drug (RLD), Prezista Tablets, 600 mg and 800 mg, of Janssen Products, L.P. The product will be launched on November 29, 2023.

The approved product has an estimated market size of US$ 274.8 million for the twelve months ending October 2023, according to IQVIA. Aurobindo now has a total of 500 ANDA approvals (478 Final approvals and 22 tentative approvals) from USFDA.

Darunavir Tablets, 600 mg and 800 mg, in combination with other antiretroviral agents, is indicated for the treatment of human immunodeficiency virus (HIV-1) infection in adult and paediatric patients 3 years of age and older.

About Aurobindo Pharma Limited

Aurobindo Pharma Limited (www.aurobindo.com), (NSE: AUROPHARMA, BSE: 524804, Reuters: ARBN.NS, Bloomberg: ARBP IN) is an integrated global pharmaceutical company headquartered in Hyderabad, India. The Company develops, manufactures, and commercializes a wide range of generic pharmaceuticals, branded specialty pharmaceuticals and active pharmaceutical ingredients globally in over 150 countries.

The company has 25 manufacturing and packaging facilities that are approved by leading regulatory agencies including USFDA, UK MHRA, EDQM, Japan PMDA, WHO, Health Canada, South Africa MCC, Brazil ANVISA. The company's robust product portfolio is spread over 7 major therapeutic/product areas encompassing CNS, Anti-Retroviral, CVS, Antibiotics, Gastroenterological, Anti-Diabetics and Anti-Allergic, supported by a strong R&D set-up.

To know more, please log on to www.aurobindo.com

For further information or queries, please contact:

Investor Relations | Corporate Communications
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Email: ir@aurobindo.com

Disclaimer:

This press release contains statements that may constitute “forward looking statements” including and without limitation, statements relating to product characteristics and uses, sales potential and target dates for product launch, implementation of strategic initiatives, and other statements relating to our future business developments and economic performance. While these forward-looking statements represent our judgment and future expectations concerning the development of our business, a number of risks, uncertainties and other factors could cause actual developments and results to differ materially from our expectations. The company undertakes no obligation to publicly revise any forward-looking statements to reflect future events or circumstances and will not be held liable for any use of this information.

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