AUROBINDO

February 19, 2021

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 Sirs,

Sub: Press Release on USFDA Approval for Droxidopa Capsules

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 Droxidopa Capsules.

Please take the information on record.

Thanking you,

Yours faithfully,
For AUROBINDO PHARMA LIMITED

AUROBINDO PHARMA LIMITED * HYDERABAD, INDIA *

B. Adi Reddy
Company Secretary

Encl: As above

AUROBINDO

NEWS RELEASE
19th Feb 2021, Hyderabad, India

Aurobindo Pharma receives USFDA Approval for Droxidopa Capsules

Aurobindo Pharma Limited is pleased to announce that the company has received final approval from the US Food & Drug Administration (USFDA) to manufacture and market Droxidopa Capsules, 100 mg, 200 mg and 300 mg. Droxidopa Capsules are generic version of Lundbeck NA Ltd's Northera® Capsules. The product will be launched immediately.

The approved product has an estimated market size of US$ 352 million for the twelve months ending December 2020, according to IQVIA. Droxidopa is indicated for the treatment of orthostatic dizziness and lightheadedness in adult patients with symptomatic neurogenic orthostatic hypotension caused by primary autonomic failure, dopamine beta-hydroxylase deficiency and non-diabetic autonomic neuropathy.

This is the 24th ANDA to be approved out of Unit-X formulation facility in Hyderabad, India used for manufacturing oral products. Aurobindo now has a total of 469 ANDA approvals (440 Final approvals and 29 tentative approvals) from USFDA.

About Aurobindo Pharma Limited:

Aurobindo Pharma Limited (www.aurobindo.com) (NSE: AUROPHARMA, BSE: 524804, Reuters: ARBN.NS, Bloomberg: ARBP:IN), headquartered at Hyderabad, India, manufactures generic pharmaceuticals and active pharmaceutical ingredients. The company's manufacturing facilities are approved by several leading regulatory agencies like US FDA, UK MHRA, Japan PMDA, WHO, Health Canada, MCC South Africa, ANVISA Brazil. The company's robust product portfolio is spread over major therapeutic/product areas encompassing Antibiotics, Anti-Retrovirals, CVS, CNS, Gastroenterologicals, Anti-Allergies and Anti-Diabetics, supported by an outstanding R&D set-up. The Company is marketing these products globally, in over 150 countries.

For further information, please contact:

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Disclaimer:

This press release contain 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.