

Date: July 30, 2018

To NATIONAL STOCK EXCHANGE OF INDIA LIMITED Exchange Plaza, Bandra Kurla Complex, Bandra (E), MUMBAI -400 051 Company Code No. AUROPHARMA	To BSE LIMITED Phiroz Jeejeebhoy Towers, 25 th floor, Dalal Street, MUMBAI -400 001 Company Code No. 524804
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Dear Sirs,

SUB: USFDA Approval for Bivalirudin Injection

We enclose copy of the Press Release issued by the Company.

Please take the information on record.

Thanking you,

Yours faithfully,
For AUROBINDO PHARMA LIMITED



B. Adi Reddy
Company Secretary

AUROBINDO PHARMA LIMITED

(CIN :L24239TG1986PLC015190)

PAN No. AABCA7366H

Corp off.: The Water Mark Building, Plot No.11, Survey No.9, Hi-tech City, Kondapur, Hyderabad – 500 084 T.S., INDIA Tel : +91 40 6672 5000 / 1200 Fax : +91 40 6707 4059

Regd. Off. : Plot No. 2, Maitrivihar, Ameerpet, Hyderabad - 500 038 T.S., INDIA Tel : +91 40 2373 6370 Fax : +91 40 2374 7340, Email : info@aurobindo.com

www.aurobindo.com

NEWS RELEASE

30 July 2018, Hyderabad, India

Aurobindo Pharma receives USFDA Approval for Bivalirudin Injection

Drug approved out of Unit IV facility

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 Bivalirudin injection, 250 mg/vial. The approved ANDA is a generic equivalent to the reference listed drug (RLD) product Angiomax® Injection of The Medicines Company.

The estimated market size of Bivalirudin injection is US\$ 101 million for the twelve months ended May 2018 according to IQVIA. The product will be launched in September 2018.

Bivalirudin injection is indicated for use as an anticoagulant in patients:

- with unstable angina undergoing percutaneous transluminal coronary angioplasty
- undergoing percutaneous coronary intervention (PCI) with provisional use of glycoprotein IIb/IIIa inhibitor (GPI) as in the REPLACE-2 study
- with, or at risk of, heparin-induced thrombocytopenia (HIT) or heparin induced thrombocytopenia and thrombosis syndrome (HITS), undergoing PCI

Bivalirudin is intended for use in these indications with aspirin.

This is the 53rd ANDA (including 1 tentative approval) to be approved out of Unit IV formulation facility in Hyderabad, India used for manufacturing general injectable & ophthalmic products. Aurobindo now has a total of 381 ANDA approvals (348 Final approvals including 17 from Aurolife Pharma LLC and 33 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 7 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.

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