

Date: June 26, 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: Press Release on USFDA Approval for Ertapenem 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

26 June 2018, Hyderabad, India

Aurobindo Pharma receives USFDA Approval for Ertapenem Injection

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 Ertapenem Injection 1 g/vial. Aurobindo's Ertapenem injection is a generic equivalent of Merck Sharp & Dohme Corp's Invanz® Injection. The product will be launched in July 2018.

The approved product has an estimated market size of US\$ 387 million for the twelve months ending Apr 2018 according to IQVIA. Ertapenem injection is used for the treatment of moderate to severe infections caused by susceptible bacteria. Also indicated in adults for the prophylaxis of surgical site infection following elective colorectal surgery.

This ANDA is approved out of Aurobindo Pharma's (wholly owned subsidiary) formulation facility in Bhiwadi, India used for manufacturing penem injectable products. Aurobindo now has a total of 371 ANDA approvals (338 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 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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