AUROBINDO

Date: April 1, 2016

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

Dear Sirs,

SUB: Press Release – Reg

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

This is for your information and record

Thanking you,

Yours faithfully,
For **AUROBINDO PHARMA LIMITED**

A.MOHAN RAMI REDDY
Vice President (Legal) & Company Secretary

AUROBINDO PHARMA LTD. * HYD. *

NEWS RELEASE
1st April 2016, Hyderabad, India

Aurobindo Pharma receives USFDA Approval for Vancomycin Hydrochloride for 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 Vancomycin Hydrochloride for Injection USP, 500 mg/vial and 1 g/vial. This product is expected to be launched in the later part of FY16-17.

The approved ANDA is bioequivalent and therapeutically equivalent to the reference listed drug product (RLD) Vancomycin® Hydrochloride for Injection USP, 500 mg/vial and 1 g/vial, of Fresenius Kabi USA, LLC.

Vancomycin Hydrochloride for Injection is an antibiotic used in the treatment of severe infections caused by susceptible strains of methicillin-resistant (Beta-lactam resistant) staphylococci and others. The approved product has an estimated market size of US$ 94 million for the twelve months ending January 2016 according to IMS.

This is the 30th ANDA (including two tentative approvals) to be approved out of Unit IV formulation facility in Hyderabad, India used for manufacturing general injectable products. Aurobindo now has a total of 250 ANDA approvals (214 Final approvals including 10 from Aurolife Pharma LLC and 36 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:
Investor Relations
Phone: 040-66725401 / 66725000
Mobile: +91 98486 67906
Email: ir@aurobindo.com

Disclaimer:
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