

Date: April 22, 2016

To <b>NATIONAL STOCK EXCHANGE OF INDIA LIMITED</b> Exchange Plaza, Bandra Kurla Complex, Bandra (E), <b>MUMBAI -400 051</b>  <b>Company Code No. AUROPHARMA</b>	To <b>BSE LIMITED</b> Phiroz Jeejeebhoy Towers, 25 <sup>th</sup> floor, Dalal Street, <b>MUMBAI -400 001</b>  <b>Company Code No. 524804</b>
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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 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](http://www.aurobindo.com)



NEWS RELEASE

22<sup>nd</sup> April 2016, Hyderabad, India

**Aurobindo Pharma receives USFDA Approval for Esomeprazole Magnesium Delayed-release 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 Esomeprazole Magnesium Delayed-release Capsules USP, 20 mg and 40 mg. The launch of this product is based on the settlement terms/litigation outcome with AstraZeneca.

The approved ANDA is bioequivalent and therapeutically equivalent to the reference listed drug product (RLD) Nexium® Delayed-release Capsules, 20 mg and 40 mg, of AstraZeneca Pharmaceuticals LP.

Esomeprazole Magnesium Delayed-release Capsules is used in the treatment of gastroesophageal reflux disease. The approved product has an estimated market size of US\$ 4.2 billion for the twelve months ending February 2016 according to IMS.

This is the 70<sup>th</sup> ANDA (including 15 tentative approvals) to be approved out of Unit VII formulation facility in Hyderabad, India for manufacturing Oral Non-Antibiotic products. Aurobindo now has a total of 253 ANDA approvals (217 Final approvals including 10 from Aurolife Pharma LLC and 36 tentative approvals) from USFDA.

**About Aurobindo Pharma Limited:**

Aurobindo Pharma Limited ([www.aurobindo.com](http://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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