

Date: January 7, 2016

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 – 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**

B. Reddy

B.ADI REDDY
Company Secretary



NEWS RELEASE

7 January 2017, Hyderabad, India

Aurobindo Pharma receives USFDA Approval for Levetiracetam in Sodium Chloride 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 Levetiracetam in Sodium Chloride Injection, 500 mg/100 mL (5 mg/mL), 1000 mg/100 mL (10 mg/mL), and 1500 mg/100 mL (15 mg/mL) (Single-use bags). Aurobindo's Levetiracetam in Sodium Chloride Injection is a generic equivalent of HQ Specialty Pharma Corp's Levetiracetam in Sodium Chloride Injection. The product will be launched in Jan 2017.

Levetiracetam in Sodium Chloride Injection is a CNS drug and indicated for partial onset seizures in adults (16 years and older) with epilepsy; myoclonic seizures in adults with juvenile myoclonic epilepsy; primary generalized tonic-clonic seizures in adults with idiopathic generalized epilepsy. The approved product has an estimated market size of US\$ 32 million for the twelve months ending November 2016 according to IMS.

This is the 41st ANDA (including 2 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 304 ANDA approvals (263 Final approvals including 16 from Aurolife Pharma LLC and 41 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:

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.

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