NEWS RELEASE
2nd February 2016, Hyderabad, India

Aurobindo Pharma receives USFDA Approval for Levetiracetam 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 Injection USP, 500 mg/5 mL (100 mg/mL) single-dose vials.

The approved ANDA is bioequivalent and therapeutically equivalent to the reference listed drug product (RLD) Keppra® Injection, 500 mg/5 mL (100 mg/mL), of UCB, Inc.

Levetiracetam Injection is used for the treatment of partial onset seizures, myoclonic seizures in patients with juvenile myoclonic epilepsy, and primary generalized tonic-clonic seizures. The approved product has an estimated market size of US$ 29 million for the twelve months ending November 2015 according to IMS.

This is the 23rd 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 236 ANDA approvals (204 Final approvals including 10 from Aurolife Pharma LLC and 32 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 6 major therapeutic/product areas encompassing Antibiotics, Anti-Retrovirals, CVS, CNS, Gastroenterologicals, and Anti-Allergies, supported by an outstanding R&D set-up. The Company is marketing these products globally, in over 150 countries.

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