

NEWS RELEASE

9<sup>th</sup> December 2015, Hyderabad, India

**Aurobindo Pharma receives USFDA Approval for Eptifibatide 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 Eptifibatide Injection USP, 20 mg/10 mL, 75 mg/100 mL, and 200 mg/100 mL single-use vials. This product is expected to be launched in this month.

The approved ANDA is bioequivalent and therapeutically equivalent to the reference listed drug product (RLD) Integrilin® Injection, 20 mg/10 mL, 75 mg/100 mL, and 200 mg/100 mL of Schering Corporation.

Eptifibatide Injection is used in the treatment of acute coronary syndrome. The approved product has an estimated market size of US\$137 million for the twelve months ending October 2015 according to IMS.

Aurobindo now has 18 ANDAs (including one tentative approval) approved out of Unit IV formulation facility in Hyderabad, India for manufacturing general injectable products. Aurobindo now has a total of 222 ANDA approvals (194 Final approvals including 10 from Aurolife Pharma LLC and 28 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 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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