

NEWS RELEASE

27th April 2016, Hyderabad, India

Aurobindo Pharma receives USFDA Approval for Famotidine Tablets (OTC)

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 Famotidine Tablets USP, 10 mg and 20 mg (OTC). This product is expected to be launched in Q2 FY16-17.

The approved ANDA is bioequivalent and therapeutically equivalent to the reference listed drug product (RLD) Pepcid® AC Tablets, 10 mg and 20 mg, of McNeil Consumer Pharmaceuticals Co.

Famotidine Tablets (OTC) is used to relieve heartburn associated with acid indigestion and sour stomach. The approved product has an estimated market size of US\$ 31 million for the twelve months ending February 2016 according to IMS.

This is the 73rd ANDA (including 16 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 256 ANDA approvals (219 Final approvals including 10 from Aurolife Pharma LLC and 37 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.

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