

NEWS RELEASE

26th April 2016, Hyderabad, India

Aurobindo Pharma receives USFDA Approval for Amlodipine and Valsartan Tablets

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 Amlodipine and Valsartan Tablets USP, 5 mg/160 mg, 10 mg/160 mg, 5 mg/320 mg and 10 mg/320 mg. This product is expected to be launched in Q1 FY16-17.

The approved ANDA is bioequivalent and therapeutically equivalent to the reference listed drug product (RLD) Exforge® Tablets of Norvartis Pharmaceuticals Corporation.

Amlodipine and Valsartan Tablets are used in the treatment of hypertension, to lower blood pressure. The approved product has an estimated market size of US\$ 123 million for the twelve months ending February 2016 according to IMS.

This is the 71st 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 254 ANDA approvals (218 Final approvals including 10 from Aurolife Pharma LLC and 36 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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