NEWS RELEASE

16th January 2016, Hyderabad, India

Aurobindo Pharma receives USFDA Approval for Paricalcitol Capsules

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 Paricalcitol Capsules, 1 mcg, 2 mcg, and 4 mcg. 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) Zemplar® capsules 1 mcg, 2 mcg and 4 mcg of ABBVIE.

Paricalcitol Capsules is used for the prevention and treatment of high levels of parathyroid hormone in certain patients with chronic kidney disease. The approved product has an estimated market size of US$ 38 million for the twelve months ending November 2015 according to IMS.

This is the 58th ANDA to be approved out of Unit VII formulation facility in Hyderabad, India for manufacturing Oral Non-Antibiotic products. Aurobindo now has a total of 231 ANDA approvals (200 Final approvals including 10 from Aurolife Pharma LLC and 31 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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