NEWS RELEASE

22nd December 2015, Hyderabad, India

Aurobindo Pharma receives USFDA Approval for Olopatadine Hydrochloride Ophthalmic Solution

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 Olopatadine Hydrochloride Ophthalmic Solution USP, 0.1%. This product is ready for launch.

The approved ANDA is bioequivalent and therapeutically equivalent to the reference listed drug product (RLD) Patanol® Ophthalmic Solution/Drops, 0.1%, of Alcon Laboratories, Inc.

Olopatadine Hydrochloride Ophthalmic Solution is used in the treatment of the signs and symptoms of seasonal allergic conjunctivitis. The approved product has an estimated market size of US$235 million for the twelve months ending October 2015 according to IMS.

This is the 20th ANDA (including one tentative approval) approved out of Unit IV formulation facility in Hyderabad, India used for manufacturing general injectable products. Aurobindo now has a total of 225 ANDA approvals (197 Final approvals including 10 from Aurolife Pharma LLC and 28 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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