NEWS RELEASE
2nd July 2018, Hyderabad, India

Aurobindo Pharma receives USFDA Approval for Ibuprofen OTC Capsules

Aurobindo Pharma Limited is pleased to announce that the Company has received final approval from the US Food & Drug Administration (USFDA) to manufacture Ibuprofen capsules OTC, 200mg. Aurobindo's Ibuprofen capsules are a generic equivalent of Pfizer's Advil Liqui-Gels Capsules. The product will be launched in Sep 2018.

Ibuprofen is used to relieve pain from various conditions such as headache, dental pain, muscular aches, and minor pain of arthritis. It is also used to reduce fever and to relieve from minor aches and pain due to the common cold or flu. The estimated market size of ibuprofen capsules OTC is US$ 164 million for the twelve months ending March 2018, according to Nielsen data.

This is the 143rd ANDA (including 19 tentative approvals) to be approved out of Unit VII formulation facility in Hyderabad, India used for manufacturing oral products. Aurobindo now has a total of 373 ANDA approvals (340 Final approvals including 17 from Aurolife Pharma LLC and 33 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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