

PRESS RELEASE

13th December 2013

Aurobindo Pharma receives USFDA approval for Duloxetine Hydrochloride Delayed Release Capsules

Aurobindo Pharma Limited is pleased to announce that the company has received the final approval from the US Food & Drug Administration to manufacture and market Duloxetine Hydrochloride Delayed-Release Capsules 20mg (base), 30mg (base) and 60mg (base) [ANDA No. 090778], which was earlier tentatively approved.

Duloxetine Hydrochloride Delayed-Release Capsules 20mg (base), 30mg (base) and 60mg (base) are the generic equivalent of Eli Lilly & Company's Cymbalta® Delayed-Release Capsules 20mg (base), 30mg (base) and 60mg (base). Duloxetine Hydrochloride Delayed-Release Capsules are indicated for the treatment of for the treatment of major depressive disorder (MDD) and falls under the Neurological (CNS) therapeutic category. The market size of the product is estimated to be US\$5.4 Billion for the twelve months ending September 2013 according to IMS.

Aurobindo now has a total of 188 ANDA approvals (163 Final approvals including 7 from Aurolife Pharma LLC and 25 Tentative approvals) from USFDA

About Aurobindo Pharma Limited:

Aurobindo Pharma Limited (www.aurobindo.com), 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, 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-Allergics, supported by an outstanding R&D set-up. The Company is marketing these products globally, in over 125 countries.

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