



Aurobindo Pharma receives tentative approval for Lopinavir/ Ritonavir Tablets

Aurobindo Pharma Limited is pleased to announce that it has received tentative approval for Lopinavir/Ritonavir Tablets 100/ 25mg and 200/ 50mg from the US Food & Drug Administration (USFDA).

Lopinavir/Ritonavir Tablets 100/ 25mg and 200/ 50mg is the generic equivalent of Abbott Laboratories' Kaletra® tablets 100/ 25mg and 200/ 50mg and falls under the Anti-Retroviral (ARV) segment. It is indicated in combination with other antiretroviral agents for the treatment of HIV-1 infection in adults and children above the age of two years. This ANDA was reviewed under the expedited provisions of the President's Emergency Plan for AIDS relief (PEPFAR).

This is the first generic product tentatively approved under PEPFAR for Abbott's Kaletra®.

Aurobindo now has a total of 93 ANDA approvals (66 Final approvals and 27 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 100 countries.

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