

NEWS RELEASE

1st April 2016, Hyderabad, India

Aurobindo Pharma receives USFDA Approval for Valganciclovir Tablets

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 Valganciclovir Tablets USP, 450 mg. This product is to be launched in Q1 FY16-17.

The approved ANDA is bioequivalent and therapeutically equivalent to the reference listed drug product (RLD) Valcyte[®] of Hoffman-La Roche Inc.

Valganciclovir Tablets is an Anti-Viral used in the treatment of Cytomegalovirus (CMV) retinitis in patients with acquired immunodeficiency syndrome and prevention of CMV disease in kidney, heart or kidney-pancreas transplant patients. The approved product has an estimated market size of US\$ 391 million for the twelve months ending February 2016 according to IMS.

This is the 69th ANDA (including 15 tentative approvals) to be approved out of Unit VII formulation facility in Hyderabad, India for manufacturing Oral Non-Antibiotic products. Aurobindo now has a total of 251 ANDA approvals (215 Final approvals including 10 from Aurolife Pharma LLC and 36 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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