

NEWS RELEASE

5th April 2016, Hyderabad, India

Aurobindo Pharma receives USFDA Approval for Polymyxin B for Injection

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 Polymyxin B for Injection USP, 500,000 units/vial. This product is expected to be launched in Q2 FY16-17.

The approved ANDA is bioequivalent and therapeutically equivalent to the reference listed drug product (RLD) Polymyxin® B for Injection USP, 500,000 units/vial of Eurohealth International Sarl.

Polymyxin B for Injection is an Anti-Infective used in the treatment of infections of the urinary tract, meninges, bloodstream and eye caused by susceptible strains of *Pseudomonas aeruginosa*. The approved product has an estimated market size of US\$ 7.6 million for the twelve months ending February 2016 according to IMS.

This is the 31st ANDA (including two tentative approvals) to be approved out of Unit IV formulation facility in Hyderabad, India used for manufacturing general injectable products. Aurobindo now has a total of 252 ANDA approvals (216 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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