

August 21, 2017

To	To
NATIONAL STOCK EXCHANGE OF INDIA LIMITED Exchange Plaza, Bandra Kurla Complex, Bandra (E), MUMBAI -400 051	BSE LIMITED Phiroz Jeejeebhoy Towers, 25 th floor, Dalal Street, MUMBAI -400 001
Company Code No. AUROPHARMA	Company Code No. 524804

Dear Sirs,

SUB: Press Release – Reg.

We enclose copy of the Press Release issued by the Company.

This is for your information and record

Thanking you,

Yours faithfully,
For **AUROBINDO PHARMA LIMITED**

B. Adi Reddy

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Company Secretary



NEWS RELEASE

21st Aug 2017, Hyderabad, India

**Aurobindo Pharma receives USFDA tentative approval for Dolutegravir, Lamivudine & Tenofovir
Disoproxil Fumarate tablets**

Tentative approval enables Aurobindo to launch the product in the PEPFAR markets

Aurobindo Pharma Limited is pleased to announce that the company has received tentative approval from the U.S. Food & Drug Administration (USFDA) under the U.S. President's Emergency Plan for AIDS Relief (PEPFAR) for its New Drug Application for Dolutegravir, Lamivudine and Tenofovir Disoproxil Fumarate (TLD) tablets, 50mg/300mg/300mg.

The approved product is used for the treatment of HIV-1 infection along as a complete regimen in adults and pediatric patients weighing 40 kg and greater. The reference listed drugs of the approved combination product are ViiV Healthcare's Tivicay® (dolutegravir) and Epivir® (Lamivudine), and Gilead Science's Viread® (Tenofovir Disoproxil Fumarate).

ViiV Healthcare and Aurobindo Pharma signed a licensing agreement in 2014 that allows Aurobindo Pharma to supply Dolutegravir 50mg in 92 licensed countries, following completion of required local regulatory approval processes.

The triple combination product is expected to be launched in sub-Saharan Africa in Q3FY18.

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.

For further information, please contact:

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Disclaimer:

This press release contain statements that may constitute "forward looking statements" including and without limitation, statements relating to product characteristics and uses, sales potential and target dates for product launch, implementation of strategic initiatives, and other statements relating to our future business developments and economic performance. While these forward looking statements represent our judgment and future expectations concerning the development of our business, a number of risks, uncertainties and other factors could cause actual developments and results to differ materially from our expectations. The company undertakes no obligation to publicly revise any forward looking statements to reflect future events or circumstances and will not be held liable for any use of this information.

AUROBINDO PHARMA LIMITED

(CIN : L24239TG1986PLC015190)

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