

May 31, 2017

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

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**  
Company Secretary



**NEWS RELEASE**

31 May 2017, Hyderabad, India

**Aurobindo Pharma receives USFDA Approval for Atomoxetine Capsules**

Aurobindo Pharma Limited is pleased to announce that the company has received final approval from the US Food & Drug Administration (USFDA) to manufacture Atomoxetine capsules, 10 mg, 18 mg, 25 mg, 40 mg, 60 mg, 80 mg and 100 mg. Atomoxetine capsules are the AB rated generic equivalent of Eli Lilly and Company's Strattera<sup>®</sup> capsules. The product will be launched immediately.

Atomoxetine capsules are indicated for the treatment of Attention-Deficit/Hyperactivity Disorder (ADHD). The approved product has an estimated market size of US\$ 1.1 billion for the twelve months ending March 2017 according to IMS.

This is the 116<sup>th</sup> ANDA (including 14 tentative approvals) to be approved out of Unit III formulation facility in Hyderabad, India used for manufacturing oral products. Aurobindo now has a total of 320 ANDA approvals (286 Final approvals including 16 from Aurolife Pharma LLC and 34 tentative approvals) from USFDA.

**About Aurobindo Pharma Limited:**

Aurobindo Pharma Limited ([www.aurobindo.com](http://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)

PAN No. AABCA7366H

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