

February 26, 2026

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

Sub: Press Release - Eugia Pharma receives USFDA Approval for Everolimus Tablets

We enclose a copy of the Press Release that is being issued by the Company in connection with USFDA approval received by Eugia Pharma Specialities Limited, a wholly owned subsidiary of the Company, for Everolimus Tablets, 0.25 mg, 0.5 mg, 0.75 mg and 1 mg.

Please take the above information on record.

Thanking you,

Yours faithfully,

For AUROBINDO PHARMA LIMITED

B. Adi Reddy
Company Secretary

Encl.: As above

(CIN : L24239TG1986PLC015190)

AUROBINDO PHARMA LIMITED
www.aurobindo.com

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Hyderabad, India, February 26, 2026

Eugia Pharma receives USFDA Approval for Everolimus Tablets

Aurobindo Pharma Limited is pleased to announce that its wholly owned subsidiary, Eugia Pharma Specialities Limited, has received final approval from the US Food & Drug Administration (USFDA) to manufacture and market Everolimus Tablets, 0.25 mg, 0.5 mg, 0.75 mg, 1 mg, which is bioequivalent and therapeutically equivalent to the reference listed drug (RLD), Zortress Tablets, 0.25 mg, 0.5 mg, 0.75 mg, 1 mg, of Novartis Pharmaceuticals Corp.

This approval has been received from Eugia Unit-I and is expected to be launched in Q1FY27.

The approved product has an estimated market size of US\$ 78 million for the twelve months ending December 2025, according to IQVIA MAT.

This is the 184th ANDA approval (including 10 tentative approvals received) out of Eugia Pharma Specialities Group (EPSG) facilities, manufacturing both oncology oral and sterile specialty products.

Everolimus Tablets are Indicated for the prophylaxis of organ rejection in adult patients (i) Kidney Transplant: at low-moderate immunologic risk. Use in combination with basiliximab, cyclosporine (reduced doses) and corticosteroids (ii) Liver Transplant: Administer no earlier than 30 days posttransplant. Use in combination with tacrolimus (reduced doses) and corticosteroids.

About Aurobindo Pharma Limited

Aurobindo Pharma Limited (www.aurobindo.com), (NSE: AUROPHARMA, BSE: 524804, Reuters: ARBN.NS, Bloomberg: ARBP IN) is an integrated global pharmaceutical company headquartered in Hyderabad, India. The Company develops, manufactures, and commercializes a wide range of generic pharmaceuticals, branded specialty pharmaceuticals and active pharmaceutical ingredients globally in over 150 countries.

The company has 30+ manufacturing and packaging facilities that are approved by leading regulatory agencies including USFDA, UK MHRA, EDQM, Japan PMDA, WHO, Health Canada, South Africa MCC, Brazil ANVISA. The Company's robust product portfolio is spread over seven major therapeutic/product areas encompassing CNS, Anti-Retroviral, CVS, Antibiotics, Gastroenterological, Anti-Diabetics and Anti-Allergic, supported by a strong R&D set-up.

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To know more, please log on to www.aurobindo.com

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Disclaimer:

This press release contains 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.

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