

March 1, 2024

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 - Aurobindo Pharma receives USFDA approval for Fingolimod Capsules, 0.5 mg.

We enclose a copy of the Press Release that is being issued by the Company in connection with USFDA approval received by the Company for Fingolimod Capsules, 0.5 mg.

Please take the information on record.

Thanking you,

Yours faithfully,
For **AUROBINDO PHARMA LIMITED**

B. Adi Reddy
Company Secretary

Encl: as above

Hyderabad, India, March 01, 2024

Aurobindo Pharma receives USFDA Approval for Fingolimod Capsules, 0.5 mg

Aurobindo Pharma Limited is pleased to announce that it has received final approval from the US Food & Drug Administration (USFDA) to manufacture and market Fingolimod Capsules, 0.5 mg, which is bioequivalent and therapeutically equivalent to the reference listed drug (RLD), Gilenya Capsules, 0.5 mg of Novartis Pharmaceuticals Corporation. The product will be launched in March 2024.

The approved product has an estimated market size of US\$ 447.3 million for the twelve months ending January 2024, according to IQVIA. Aurobindo now has a total of 505 ANDA approvals (486 Final approvals and 19 tentative approvals) from USFDA.

Fingolimod Capsules, 0.5 mg, is indicated for the treatment of patients with relapsing forms of multiple sclerosis (MS) to reduce the frequency of clinical exacerbations and to delay the accumulation of physical disability.

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 25 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 7 major therapeutic/product areas encompassing CNS, Anti-Retroviral, CVS, Antibiotics, Gastroenterological, Anti-Diabetics and Anti-Allergic, supported by a strong R&D set-up.

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

For further information or queries, please contact:

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AUROBINDO PHARMA LIMITED

(CIN : L24239TG1986PLC015190)

www.aurobindo.com

PAN No. AABCA7366H

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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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