

June 4, 2026

To Listing Department, NATIONAL STOCK EXCHANGE OF INDIA LIMITED Exchange Plaza, Bandra Kurla Complex, Bandra (E), MUMBAI -400 051 Company Code No. AUOPHARMA	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 final US FDA approval for Tofacitinib Tablets, 5 mg and 10 mg

We enclose a copy of the Press Release that is being issued by the Company in connection with receipt of final US FDA approval for Tofacitinib Tablets, 5 mg and 10 mg.

Please take the information on record.

Thanking you,

Yours faithfully,
For **AUROBINDO PHARMA LIMITED**

B. Adi Reddy
Company Secretary

Encl: as above

AUROBINDO PHARMA LIMITED

www.aurobindo.com

(CIN : L24239TG1986PLC015190)

Corp. Off.: Galaxy, Floors: 22-24, Plot No.1, Survey No.83/1, Hyderabad Knowledge City, Raidurg Panmaktha, Ranga Reddy District, Hyderabad – 500 032, Telangana, India.

Tel : +91 40 6672 5000 / 6672 1200 Fax: +91 40 6707 4044.

Regd. off.: Plot No. 2, Maithrivihar, Ameerpet, Hyderabad - 500 038, Telangana., India. Tel: +91 40 2373 6370/ 2374 7340 Fax: +91 40 2374 1080 / 2374 6833

Email: info@aurobindo.com Website: www.aurobindo.com

Hyderabad, India, June 4, 2026

Aurobindo Pharma receives final US FDA approval for Tofacitinib Tablets, 5 mg and 10 mg

Aurobindo Pharma Limited is pleased to announce that it has received a final approval from the US Food & Drug Administration (USFDA) to manufacture and market Tofacitinib Tablets, 5 mg and 10 mg, which is bioequivalent and therapeutically equivalent to the reference listed drug (RLD), Xeljanz Tablets, 5 mg and 10 mg, of PF Prism C.V.

The product will be manufactured at APL Healthcare Unit IV, a wholly owned subsidiary of the company and will be launched immediately.

According to IQVIA MAT data for the twelve months ending April 2026, Tofacitinib Tablets, 5 mg and 10 mg have an estimated market size of approximately US\$ 494 million in the U.S.

Aurobindo Pharma now has a total of 586 ANDA approvals (561 final approvals and 25 tentative approvals) from USFDA.

Tofacitinib Tablets are indicated for: (1) the treatment of adult patients with moderately to severely active rheumatoid arthritis who have had an inadequate response or intolerance to methotrexate, (2) the treatment of adult patients with active psoriatic arthritis who have had an inadequate response or intolerance to methotrexate or other disease-modifying antirheumatic drugs (DMARDs), (3) the treatment of adult patients with moderately to severely active ulcerative colitis (UC), who have had an inadequate response or who are intolerant to TNF blockers.

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.

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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