

April 7, 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 - Aurobindo Pharma receives USFDA approval for Dapagliflozin and Metformin Hydrochloride Extended-Release Tablets

We enclose a copy of the Press Release that is being issued by the Company in connection receipt of USFDA approval for Dapagliflozin and Metformin Hydrochloride Extended-Release Tablets, 5 mg/ 500 mg, 5 mg/1000 mg, 10 mg/500 mg, and 10 mg/1000 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, April 07, 2026

Aurobindo Pharma receives USFDA Approval for Dapagliflozin and Metformin Hydrochloride Extended-Release Tablets, 5 mg/ 500 mg, 5 mg/1000 mg, 10 mg/500 mg, and 10 mg/1000 mg

Aurobindo Pharma Limited is pleased to announce that the final approval is received from the US Food & Drug Administration (USFDA) to manufacture and market Dapagliflozin and Metformin Hydrochloride Extended-Release Tablets, 5 mg/ 500 mg, 5 mg/1000 mg, 10 mg/500 mg, and 10 mg/1000 mg, which is bioequivalent and therapeutically equivalent to the reference listed drug (RLD), Xigduo XR Tablets, 5 mg/ 500 mg, 5 mg/1000 mg, 10 mg/500 mg, and 10 mg/1000 mg, of AstraZeneca AB.

These products will be manufactured at Unit-IV of APL Healthcare Limited, a wholly owned subsidiary of the Company and will be launched immediately.

The approved product has an estimated market size of US\$ 514 Million for the twelve months ending February 2026, according to IQVIA MAT. Aurobindo Pharma has a total of 579 ANDA approvals (554 final approvals and 25 tentative approvals) from USFDA as of 31 March 2026.

Aurobindo Pharma Limited, being one of the first ANDA applicants to submit a substantially complete ANDA with a paragraph IV certification for Dapagliflozin and Metformin Hydrochloride Extended-Release Tablets, 5 mg/ 500 mg, 5 mg/1000 mg, 10 mg/500 mg, and 10 mg/1000 mg, is thus eligible for 180 days of shared generic drug exclusivity.

Dapagliflozin and Metformin Hydrochloride Extended-Release Tablets are indicated as an adjunct to diet and exercise to improve glycaemic control in adults with type-2 diabetes mellitus when treatment with both dapagliflozin and metformin is appropriate.

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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