

May 13, 2025

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 - CuraTeQ Biologics receives approval for biosimilar Zefylti from UK's MHRA

We enclose a copy of the Press Release that is being issued by the Company in connection with the receipt of approval for biosimilar Zefylti from UK's MHRA by CuraTeQ Biologics s.r.o., a wholly owned step-down subsidiary of the Company.

Please take the 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

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, Maithriviham, 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, May 13, 2025

CuraTeQ Biologics receives approval for biosimilar Zefylti from UK's MHRA

CuraTeQ Biologics s.r.o., a wholly owned step-down subsidiary of Aurobindo Pharma Ltd, has obtained marketing authorisation from UK's Medicines and Healthcare products Regulatory Agency (MHRA) for Zefylti®, its filgrastim biosimilar version. Earlier in February 2025, Zefylti® has received the marketing authorisation in the European Union from the European Commission (EC). This is CuraTeQ's second biosimilar to be approved by MHRA after the approval of Bevqolva® in December 2024.

About CuraTeQ Biologics Private Limited

CuraTeQ Biologics Private Limited (CuraTeQ), a wholly owned subsidiary of Aurobindo Pharma Limited, is a global biopharmaceutical company headquartered in Hyderabad, India. CuraTeQ's vision is to improve the wellbeing of patients suffering from debilitating illnesses by providing them access to high quality and cost-effective biosimilars. It is focused on developing biosimilars for the treatment of various cancers and autoimmune diseases. CuraTeQ's pipeline consists of fourteen biosimilars, primarily targeting the immunology and oncology segments. It has end-to-end capabilities in producing a full range of products from bulk drug substance to fill-finish and packaged drug products.

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

For further information or queries, please contact:

Investor Relations
Phone: +91 40 66721551 / 66725000
Email: ir@aurobindo.com

Corporate Communications
Phone: +91 40 66725005 / 66725000
Email: ccommunications@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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