

December 13, 2024

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

**Sub: Press Release - CuraTeQ Biologics receives positive opinion for biosimilar Zefylti**

We enclose a copy of the Press Release that is being issued by the Company announcing that the Committee for Medicinal Products for Human Use (CHMP) within the European Medicines Agency (EMA) has adopted a positive opinion for Zefylti (BP13, a filgrastim biosimilar) recommending the granting of a marketing authorisation to CuraTeQ Biologics Private Limited, a wholly owned subsidiary of the Company.

Please take the information on record.

Thanking you,

Yours faithfully,  
For **AUROBINDO PHARMA LIMITED**

B. Adi Reddy  
Company Secretary

Encl.: Annexure

Hyderabad, India, December 13, 2024

### CuraTeQ Biologics receives positive opinion for biosimilar Zefylti

CuraTeQ Biologics Private Limited, a wholly owned subsidiary of Aurobindo Pharma Limited, is pleased to announce that the Committee for Medicinal Products for Human Use (CHMP) within the European Medicines Agency (EMA) has adopted a positive opinion for Zefylti (BP13, a filgrastim biosimilar) recommending the granting of a marketing authorisation.

Zefylti will be available as 30 MU/0.5 mL and 48 MU/0.5 mL and is intended for the treatment of neutropenia and the mobilisation of peripheral progenitor cells (PBPCs).

Commenting on the update the company spokesperson said, "After receiving the GMP certificate of compliance from the European Medicines Agency (EMA) in November, Zefylti is our first biosimilar to receive a positive opinion from EMA's CHMP. Two more products, our biosimilar versions of pegylated filgrastim (BP14) and trastuzumab (BP02), are currently under review with the EMA, and a biosimilar to bevacizumab (BP01) is under review with the MHRA. We are confident and on track to bring these treatment options to patients next year".

### About CuraTeQ Biologics Private Limited

CuraTeQ Biologics Private Limited (CuraTeQ), a 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](http://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 29 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](http://www.aurobindo.com)

## AUROBINDO PHARMA LIMITED

(CIN : L24239TG1986PLC015190)

[www.aurobindo.com](http://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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