April 26, 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, 25th floor, Dalal Street, **MUMBAI -400 001** **Company Code No. 524804**

Dear Sir/ Madam,

Sub: Press Release - CuraTeQ Biologics s.r.o, a wholly owned step-down subsidiary of the Company, receives Positive Opinion from CHMP for Dazublys®, a Trastuzumab Biosimilar

We enclose a copy of the Press Release that is being issued by CuraTeQ Biologics s.r.o, a wholly owned step-down subsidiary of the Company in connection with the receipt of Positive Opinion from CHMP for Dazublys®, a Trastuzumab Biosimilar.

Please take the information on record.

Thanking you,

Yours faithfully,
For **AUROBINDO PHARMA LIMITED**

Digitally signed by BADDIGAM ADI REDDY
Date: 2025.04.26 09:21:55 +05'30'

B. Adi Reddy
Company Secretary

Encl: as above

Hyderabad, India, April 26, 2025

CuraTeQ Biologics receives positive opinion for biosimilar Dazublys® from EMA

CuraTeQ Biologics s.r.o., a wholly owned step-down subsidiary of Aurobindo Pharma Limited, is pleased to announce that the Committee for Medicinal Products for Human Use (CHMP) of the European Medicines Agency (EMA) has adopted a positive opinion recommending marketing authorisation of Dazublys® (150 mg powder for concentrate for solution for infusion), its trastuzumab biosimilar, for the treatment of HER2-positive metastatic and early breast cancers.

Trastuzumab specifically binds and inhibits the human epidermal growth factor receptor 2 (HER2) protein, which is over-expressed on certain types of solid cancers such as breast and gastric cancer. By binding to the extracellular domain of HER2, trastuzumab disrupts its ability to signal, leading to cell cycle arrest, reduced tumor growth, and potentially immune system activation to destroy cancer cells.

Commenting on the update, Dr. Satakarni Makkapati, Director of Aurobindo Pharma and CEO Biologics, Vaccines and Peptides stated, "The CHMP's positive opinion is based on demonstrating comprehensive analytical similarity and clinically no meaningful differences between Dazublys® and the reference biologic product Herceptin® in terms of pharmacokinetics (PK), pharmacodynamics (PD), efficacy, safety, and immunogenicity. Upon European Commission approval that is expected in July, Dazublys® will be available for use across EU member states. This marks our third biosimilar to receive CHMP's endorsement and the fourth overall in the EU, alongside the approval of Bevqolva® (a bevacizumab biosimilar) by the MHRA in November 2024. Biosimilars are playing an important role in improving cancer care, and we remain committed to expanding our biosimilars portfolio to address the unmet needs of patients."

Aurobindo Pharma's Vice Chairman and Managing Director Mr. Nithyananda Reddy said, "The CHMP's positive opinion of a third biosimilar from our portfolio in a five-month time period underscores our extensive efforts in building biosimilars as one of the core businesses at Aurobindo. By 2030, we are committed to launching at least 10 biosimilars across oncology and immunology therapy segments, furthering our mission to serve patients with high-quality, cost-effective therapies."

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), (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:

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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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