March 15, 2024

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 announces the successful Phase 1 clinical study outcome of their BP11 product (a biosimilar to Xolair)

We enclose a copy of the Press Release that is being issued by the Company in connection with announcement by CuraTeQ Biologics Private Limited, a wholly owned subsidiary of the Company, the successful Phase 1 clinical study outcome of their BP11 product (a biosimilar to Xolair).

Please take the information on record.

Thanking you,

Yours faithfully,
For **AUROBINDO PHARMA LIMITED**

BADDIGAM ADI REDDY
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B. Adi Reddy
Company Secretary

Encl: as above

Hyderabad, India, March 15, 2024

CuraTeQ Biologics announces the successful Phase 1 clinical study outcome of their BP11 product (a biosimilar to Xolair)

CuraTeQ Biologics Private Limited, a wholly owned subsidiary of Aurobindo Pharma Ltd, announced that their Omalizumab biosimilar candidate BP11 has met the Phase 1 trial end points vis-à-vis the EU and US sourced reference product Xolair. The PK/PD trial was conducted in 165 healthy volunteers in Australia and New Zealand.

"The primary objective was to prove pharmacokinetic (PK) equivalence between BP11, US and EU sourced Xolair. 165 healthy volunteers were randomized to receive either BP11 or EU or US licensed Omalizumab via subcutaneous route of administration. Results of both primary parameters, i.e. maximum serum concentration (C_{max}) and area under concentration-time curve from time zero to infinity ($AUC_{0\text{-}inf}$), were contained within 80-125% bioequivalence limit demonstrating PK equivalence between BP11 and both US and EU sourced Xolair. BP11 also had similar IgE levels to Xolair demonstrating comparable pharmacodynamic profile versus US and EU sourced Xolair. The safety and immunogenicity profiles were also found comparable versus the originator's product," said Dr. Arpit Prajapati, Head of Clinical Sciences at CuraTeQ Biologics.

Dr. Disha Dadke, Associate President and Head R&D, said, "We have initiated a Phase 3 study of our Omalizumab candidate BP11 for the treatment of chronic spontaneous or idiopathic urticaria, which is a presence of hives that are itchy and can last for a number of weeks with no apparent external trigger. The Phase 3 efficacy and safety study is being conducted across multiple sites in seven European countries and in 600 patients with chronic spontaneous urticaria. Additionally, a separate Phase 3 trial in asthma patients is being carried out in the Indian population. CuraTeQ intends to file the Omalizumab biosimilar product in India in 2024 and is on track to file the product in regulated markets in 2025."

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 25 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 7 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:

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

Kavita Jayaram
Corporate Communications
Phone: +91 40 66725005 / 66725000
Email: ccommunications@aurobindo.com

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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