

July 21, 2026

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

Sub: ANVISA Approval for CuraTeQ Biologics' Biosimilars Manufacturing Facility

Pursuant to Regulation 30 of the SEBI (Listing Obligations & Disclosure Requirements) Regulations, 2015, this is to inform you that:

CuraTeQ Biologics Private Limited, a wholly owned subsidiary of Aurobindo Pharma Limited, has received approval from ANVISA (Agência Nacional de Vigilância Sanitária), the National Health Surveillance Agency of Brazil, for its biosimilars manufacturing facility in Hyderabad.

The facility underwent a GMP inspection by ANVISA from May 11 to May 15, 2026, and the approval was granted following successful completion of the inspection for biological API and sterile products. The scope of the GMP inspection covered four biosimilar products and assessed the mammalian and microbial drug substance manufacturing facility, prefillable syringe and vial filling facility, packaging and labelling facility, QC testing and release laboratories.

CuraTeQ's drug substance and drug product GMP facility is already EMA (European Medicines Agency) and WHO certified and the ANVISA approval reaffirms CuraTeQ Biologics' commitment to global regulatory compliance and quality standards. The approval strengthens the Company's capability to supply biosimilar products to the Brazilian and broader Latin American markets.

Please take the above information on record.

Yours faithfully,

For AUROBINDO PHARMA LIMITED

B. Adi Reddy
Company Secretary

AUROBINDO PHARMA LIMITED

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(CIN : L24239TG1986PLC015190)

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