

January 9, 2026

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: CuraTeQ Biologics receives NOC from Health Canada for its biosimilar Dyrupreg

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

CuraTeQ Biologics Private Limited, a wholly owned subsidiary of Aurobindo Pharma Ltd, has obtained Notice of Compliance (NOC) from Biologics and Radiopharmaceutical Drugs Directorate (BRDD) for Dyrupreg™, its pegylated filgrastim biosimilar version. A Notice of Compliance (NOC) from Health Canada is issued to a drug manufacturer after a successful review, confirming the product meets regulatory standards for safety, efficacy, and quality under the Food and Drug Regulations. For Dyrupreg™, the NOC specifically indicates that Health Canada has verified high similarity to an approved reference biologic drug, with no clinically meaningful differences in terms of safety, PK/PD, or quality attributes. In 2025, Dyrupreg™ received marketing authorization in the European Union from the European Commission (EC) and from MHRA, UK. There are three other biosimilar applications from CuraTeQ Biologics seeking marketing authorisation that are currently under review with Health Canada.

Please take the information on record.

Yours faithfully,
For AUROBINDO PHARMA LIMITED

B. Adi Reddy
Company Secretary

AUROBINDO PHARMA LIMITED

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