

CONCORD BIOTECH LIMITED

B-1601-1602, B-wing Mondeal Heights, Iskcon Cross Road, S. G. Highway, Ahmedabad-380015, Gujarat.

Phone : +91-79-68138700 Fax : +91-79-68138725 CIN No.: L24230GJ1984PLC007440

Email ID: complianceofficer@concordbiotech.com

June 10, 2026

To The Manager, Listing Department National Stock Exchange of India Limited Plot No. C/1 G Block, Bandra-Kurla Complex, Bandra (East), Mumbai -400 051 Symbol: CONCORDBIO	To General Manager, Listing Department BSE Limited Phiroze Jeejeebhoy Towers, Dalal Street, Mumbai – 400 001 Scrip Code: 543960
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Sub.: Concord Biotech Secures Abbreviated New Drug Application (ANDA) Approval for Tofacitinib Tablets, 5 mg and 10 mg

Dear Sir/Madam,

Pursuant to Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015, we are pleased to inform that the Company has received approval from the U.S. Food and Drug Administration (USFDA) for its ANDA for Tofacitinib Tablets, 5 mg and 10 mg. Tofacitinib tablets are indicated for the treatment of adult patients with Moderately to severely active rheumatoid arthritis (RA), Active psoriatic arthritis (PsA), Active ankylosing spondylitis (AS), Moderately to severely active ulcerative colitis (UC), Active PsA, Active polyarticular course juvenile idiopathic arthritis (pcJIA).

According to market estimates, the U.S. market for Tofacitinib Tablets both strengths put together is approximately US\$ 500 million. The approval positions the Company to capitalize on these attractive market opportunities, enhance its product offerings, and support its long-term growth strategy across the U.S. and international markets.

The requisite details of terms of the provisions of Regulation 30 read with Schedule III and SEBI Master Circular no. SEBI/HO/CFD/PoD2/CIR/P/0155 dated November 11, 2024 read with Master Circular no. HO/49/14/14(7)2025-CFD-POD2/I/3762/2026 issued on July 11, 2023 and last updated on January 30, 2026 are enclosed herewith.

This is for your information and records.

For Concord Biotech Limited

Paritosh Trivedi
Company Secretary & Compliance Officer
ACS 63623

Encl : As above

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Sr. No.	Particulars	Details
a)	name of the regulatory or licensing authority	U. S. Food & Drug Administration (USFDA)
b)	brief details of the approval/license obtained/ withdrawn/ surrendered	Approval for Tofacitinib Tablets, 5 mg and 10 mg,
c)	impact/relevance of such approval/license to the listed entity	The approval for Tofacitinib Tablets, 5 mg and 10 mg is in line with Concord Biotech's growth strategy and will strengthen Company's position in the U.S. market.
d)	withdrawal/cancellation or suspension of licence/approval by the regulatory or licensing authority, with reasons for such action, estimated impact (monetary or otherwise) on the listed entity and penalty, if any	Not Applicable
e)	period for which such approval/license is/was valid	Not Applicable
f)	the actual impact (monetary or otherwise) along with corrective actions taken by the listed entity pursuant to the withdrawal, cancellation or suspension of the key license/ approval.	Not Applicable