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CIN: L24234KA1978PLC003417

www.biocon.com

BIO/SECL/SG/2024-25/134

December 01, 2024

То	То
The Manager,	The Manager,
BSE Limited	National Stock Exchange of India Limited
Department of Corporate Services	Corporate Communication Department
Phiroze Jeejeebhoy Towers,	Exchange Plaza, Bandra Kurla Complex
Dalal Street, Mumbai – 400 001	Mumbai – 400 050
Scrip Code - 532523	Scrip Symbol - Biocon

Dear Sir/Madam,

Subject: Notification to Stock Exchanges

Please find enclosed the company statement titled "U.S.FDA Approves Biocon Biologics' YESINTEK™, Bmab 1200 biosimilar to J&J's Stelara® (Ustekinumab)".

The above information will also be available on the website of the Company at www.biocon.com.

Kindly take the same on record and acknowledge.

Thanking You,

Yours faithfully,

For **Biocon Limited**

Mayank Verma

Meinel.

Company Secretary & Compliance Officer

Membership No: ACS 18776

Encl: as above



NOTIFICATION TO STOCK EXCHANGE

COMPANY STATEMENT

U.S.FDA Approves Biocon Biologics' YESINTEK™, Bmab 1200 biosimilar to J&J's Stelara® (Ustekinumab)

Bengaluru, Karnataka, India, December 01, 2024

Biocon Biologics Ltd (BBL), a fully integrated global biosimilars company and subsidiary of Biocon Ltd (BSE code: 532523, NSE: BIOCON), announced today that the **U.S. Food and Drug Administration (FDA) has approved YESINTEK™** (*Ustekinumab-kfce*), a biosimilar to the reference product, Stelara® (*Ustekinumab*).

YESINTEK™, a monoclonal antibody, is approved for the treatment of Crohn's disease, Ulcerative Colitis, Plaque Psoriasis and Psoriatic Arthritis.

Biocon Biologics Ltd had previously notified the Stock Exchange on Feb 29, 2024, that the Company had entered into a settlement and licensing agreement with Janssen Biotech Inc., Janssen Sciences Ireland, and Johnson & Johnson (collectively known as Janssen) to commercialize YESINTEK™ in the United States of America no later than on February 22, 2025, upon approval from the U.S. FDA.

- Company Spokesperson

For more information: seema.ahuja@biocon.com