

11th April 2025

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| <p>(1) BSE Limited
Listing Department,
Phiroze Jeejeebhoy Towers,
Dalal Street,
Mumbai 400 001
Scrip Code: 500087</p> | <p>(2) National Stock Exchange of India Limited
Listing Department,
Exchange Plaza, 5th floor,
Plot no. C/1, G Block,
Bandra Kurla Complex,
Bandra (East), Mumbai - 400 051
Scrip Code: CIPLA EQ</p> |
| <p>(3) SOCIETE DE LA BOURSE DE LUXEMBOURG
Societe Anonyme
35A Boulevard Joseph II,
L-1840 Luxembourg</p> | |

Dear Sir / Madam,

Sub: Intimation under Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015

Pursuant to the provisions of Regulation 30 read with Schedule III Part A Para B of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 and the SEBI Circular no. SEBI/HO/CFD/PoD2/CIR/P/0155 dated 11th November, 2024, we hereby notify that the Company has received final approval from the United States Food and Drug Administration ('USFDA') for the Abbreviated New Drug Application ('ANDA') submitted for Paclitaxel Protein-bound Particles for Injectable Suspension (albumin-bound), 100 mg/vial, Single-Dose Vial ('Protein-bound Paclitaxel') on 10th April, 2025.

Cipla's Protein-bound Paclitaxel is an AB-rated generic therapeutic equivalent version of Bristol Myers Squibb's Abraxane® for Injectable Suspension 100 mg/vial. Protein-bound Paclitaxel is indicated for the treatment of metastatic breast cancer, locally advanced or metastatic non-small cell lung cancer ('NSCLC') and metastatic adenocarcinoma of the pancreas. The product is expected to be launched in H1 FY 2025-26 in the United States of America.

Yours faithfully,
For Cipla Limited

Rajendra Chopra
Company Secretary

Prepared by: Mandar Kurghode