

30th May 2025

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| <p>(1) BSE Limited
Listing Department,
Phiroze Jeejeebhoy Towers,
Dalal Street,
Mumbai 400 001</p> <p>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</p> <p>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: USFDA inspection at Company's manufacturing facility in Bommasandra, Bengaluru, India

Pursuant to Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015, we hereby notify that United States Food and Drugs Administration ('USFDA') has conducted a current Good Manufacturing Practices ('cGMP') inspection at the Company's manufacturing facility in Bommasandra, Bengaluru from 26th to 30th May 2025.

On conclusion of the inspection, the Company received 1 (one) observation in Form 483. The Company will work closely with the USFDA and remains committed to address the observation comprehensively within stipulated time.

Please take the above information on record.

Yours faithfully,
For Cipla Limited

Rajendra Chopra
Company Secretary

Prepared by: Mandar Kurghode