

21<sup>st</sup> June 2024

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

Dear Sir/Madam,

**Sub: USFDA inspection at Company's manufacturing facility in Goa, India**

Pursuant to Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 and further to our earlier intimations on the captioned subject dated 28<sup>th</sup> September 2019, 22<sup>nd</sup> January 2020, 26<sup>th</sup> February 2020, 26<sup>th</sup> August 2022 and 24<sup>th</sup> November 2022, we hereby notify that the USFDA has conducted an inspection at the Company's manufacturing facility in Goa, India from 10<sup>th</sup> – 21<sup>st</sup> June 2024.

On conclusion of the inspection, the Company received 6 (six) inspectional observations in Form 483. The Company will work closely with the USFDA and is committed to address these comprehensively within stipulated time.

Please take the above information on record.

Yours faithfully,  
**For Cipla Limited**

**Rajendra Chopra**  
**Company Secretary**

Prepared by: Mandar Kurghode