

18<sup>th</sup> November, 2023

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| <p>(1) BSE Ltd<br/>Listing Department<br/>Phiroze Jeejeebhoy Towers,<br/>Dalal Street,<br/>Mumbai 400 001<br/><b>Scrip Code: 500087</b></p> | <p>(2) National Stock Exchange of India Ltd<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<br/><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: Update on USFDA inspection at Cipla's manufacturing facility in Pithampur, Indore**

Pursuant to Regulation 30 of SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 and in furtherance to our intimation dated 18<sup>th</sup> February 2023 and 5<sup>th</sup> August 2023, we wish to inform you that on 18<sup>th</sup> November, 2023, the Company has received a Warning Letter dated 17<sup>th</sup> November, 2023 from United States Food and Drug Administration (USFDA) for the routine current Good Manufacturing Practices (cGMP) inspection conducted at our Pithampur manufacturing facility between 6<sup>th</sup> – 17<sup>th</sup> February, 2023. This Warning Letter summarizes contraventions regarding methods or controls followed at the facility which do not conform to the prescribed cGMP regulations and contains directional guidance for necessary corrections.

The Company will respond to the Warning Letter within the stipulated timelines and work closely with the USFDA to address the concerns in a holistic and timely manner to ensure sustained compliance. We uphold quality and compliance with utmost importance and remain committed to be compliant with the cGMP quality standards.

Please take the above information on record.

Yours faithfully,  
**For Cipla Limited**

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

Prepared by: Mandar Kurghode