

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

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| (1) BSE Limited<br>Listing Department,<br>Phiroze Jeejeebhoy Towers,<br>Dalal Street,<br>Mumbai 400 001<br><br><b>Scrip Code: 500087</b> | (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><br><b>Scrip Code: CIPLA</b> |
| (3) SOCIETE DE LA BOURSE DE<br>LUXEMBOURG<br>Societe Anonyme<br>35A Boulevard Joseph II,<br>L-1840 Luxembourg                            |   |

**Sub: USFDA cGMP inspection completed at Cipla's manufacturing facility in Pithampur, Indore**

Dear Sir / Madam,

Pursuant to Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015, we hereby notify that the United States Food and Drug Administration (USFDA) conducted a current Good Manufacturing Practices (cGMP) inspection at our Pithampur manufacturing facility from 6<sup>th</sup> – 17<sup>th</sup> February, 2023.

On conclusion of the inspection, the Company has received 8 inspectional observations in Form 483. The Company will work closely with the USFDA and is committed to address these comprehensively within stipulated time.

Kindly take the above information on record.

Thanking you

Yours faithfully,  
**For Cipla Limited**

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

Prepared by: Mandar Kurghode