

24<sup>th</sup> November, 2022

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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>                                |                                                                                                                                                                                                                                                  |

**Sub: Update on USFDA inspection at Cipla's manufacturing facility in Goa**

Dear Sir/Madam,

Pursuant to Regulation 30 of SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 and in furtherance to intimation dated 26<sup>th</sup> August, 2022 for United States Food and Drug Administration (USFDA) inspection of our Goa manufacturing facility, we wish to inform you that the Company has received a communication from the USFDA that the classification of Company's said facility continues to be as Official Action Indicated ("OAI"). USFDA may continue to withhold product approvals from this facility till the outstanding observations are resolved. The Company has an ongoing derisking plan in place for new product approvals. The Company will work closely with the USFDA and is committed to address these within the stipulated time.

Kindly take the above information on record.

Thanking you,  
Yours faithfully,  
**For Cipla Limited**

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

Prepared by: Mandar Kurghode