

16<sup>th</sup> April 2018

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| <p>(1) BSE Ltd<br/>Listing Department<br/>Phiroze Jeejeebhoy Towers,<br/>Dalal Street,<br/>Mumbai - 400 001</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</p> |
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**Scrip Code: 500087**

**Scrip Code: CIPLA EQ**

- (3) SOCIETE DE LA BOURSE DE LUXEMBOURG  
Societe Anonyme  
35A Boulevard Joseph II,  
L-1840 Luxembourg

Dear Sirs,

The Company wishes to inform you that the US FDA conducted a routine current Good Manufacturing Practices (cGMP) audit at its Indore formulations facility from 2<sup>nd</sup> April 2018 till 13<sup>th</sup> April 2018. This was in the normal course of business. There were no data integrity and/or repeat observations. At this stage, the Company believes the observations are unlikely to have any material adverse impact. The Company will submit the response within the stipulated timeframe.

Thank you,

Yours faithfully,  
For Cipla Limited



Kedar Upadhye  
Global Chief Financial Officer