

20th July, 2019

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| <p>(1) BSE Ltd
Listing Department
Phiroze Jeejeebhoy Towers,
Dalal Street,
Mumbai - 400 001</p> | <p>(2) National Stock Exchange of India Ltd
Listing Department
Exchange Plaza, 5th floor,
Plot no. C/1, G Block,
Bandra Kurla Complex,
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

Subject: USFDA cGMP inspection completed at Cipla's API manufacturing facility in Virgonagar, Bengaluru

This is further to our letter dated 19th July 2019 regarding the captioned subject. The dates of inspection were inadvertently mentioned as 15th June 2019 to 19th June 2019 instead of 15th July 2019 to 19th July 2019. Please read the revised intimation as under:

"Pursuant to Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015, we would like to inform you that the United States Food and Drug Administration (USFDA) conducted a routine cGMP inspection at our API manufacturing facility in Virgonagar, Bengaluru, from 15th July, 2019 to 19th July, 2019.

The inspection ended with 7 observations, none of which were repeat or related to data integrity. The Company will respond to the agency within the stipulated timeline."

Kindly take the above information on record.

Thanking you,

Yours faithfully,

For **Cipla Limited**



Karan Tanna
Associate Director
Corporate Secretarial