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CIN : L24234KA1978PLC003417

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Date of submission:December 17, 2018

To The Secretary BSE Limited Department of Corporate Services PhirozeJeejeebhoy Towers, Dalal Street, Mumbai – 400 001 Scrip Code - 532523	To The Secretary National Stock Exchange of India Limited Exchange Plaza, BandraKurla Complex Mumbai – 400 050 Scrip Code- BIOCON

Dear Sir/Madam,

Sub: Biocon's APIs Manufacturing Facility in Telangana Completes U.S. FDA Inspection with No 483 Observations

Ref: Regulation 30 of SEBI Listing Obligations and Disclosure Requirements (LODR) Regulations, 2015

Pursuant to Regulation 30 of the SEBI LODR Regulations, 2015, please find below the "Company Statement" on the subject matter.

"This is to inform you that the U.S. FDA conducted a GMP inspection of our APIs manufacturing facility at Telangana from Dec 12- Dec 14, 2018. The inspection concluded without any observations and no Form 483 was issued. The successful inspection of this site reflects our strong commitment to quality and cGMP compliance." - Company Spokesperson

We request you to kindly take this to your records as per the requirement of LODR and oblige.

Thanking You,
Yours faithfully
For Biocon Limited

Satish Kumar S S
Company Secretary and Compliance Officer