



**Biocon Limited**  
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Date of submission: March 6, 2019

To The Secretary BSE Limited Department of Corporate Services Phiroze Jeejeebhoy Towers, Dalal Street, Mumbai – 400 001 Scrip Code - 532523	To The Secretary National Stock Exchange of India Limited Exchange Plaza, Bandra Kurla Complex Mumbai – 400 050 Scrip Code- BIOCON
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Dear Sir/Madam,

**Sub: Biocon Facility Completes Pre Approval U.S. FDA Inspection**

**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 concluded a pre-approval inspection of Biocon's insulin drug substance manufacturing facility triggered by a New Drug Application submitted by our insulin API customer. The inspection at the Bengaluru facility took place between 25th Feb – 5th Mar, 2019, resulting in a Form 483 with six observations. Biocon is confident of addressing these expeditiously and remains committed to global standards of Quality and 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 & Compliance Officer