

**Biocon Limited** 

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www.biocon.com

June 22, 2022
To,
The Secretary **BSE Limited**Department of Corporate Services,
Phiroze Jeejeebhoy Towers,
Dalal Street, Mumbai – 400 001

Dear Sir/Madam,

**Scrip Code: 532523** 

Subject: Clarification on news item appearing in media/publication

This is with reference to your email dated June 21, 2022, seeking clarification with regards to the media report titled, "Biocon subsidiary caught offering bribe to waive off trial of new drug".

In this regard, please note that the Company vide its letter dated June 21, 2022, has submitted the Company Statement issued by Biocon Biologics, to the stock exchanges. The aforesaid letter is attached herewith for reference.

The Company has always ensured that all material information is disclosed under Regulation 30 of the SEBI (Listing Obligation and Disclosure Requirements) Regulations, 2015 ('SEBI Listing Regulations').

We confirm that there is no unpublished price sensitive information available with the Company and there is no information that is material and not disclosed to the stock exchanges. We are cautious of the regulatory provisions and will be informing the stock exchanges of any such event as per the materiality policy of our company.

We request you to kindly take the above clarification on record.

Thanking you,

Yours faithfully,

For Biocon Limited

Mayank Verma

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Company Secretary and Compliance Officer



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June 21, 2022

Bengaluru, India

Biocon Biologics issues a statement in response to the media reports related to bribery allegations.

We strongly deny the allegations of bribery against the Company and its officials associated with the approval process of one of our products in India.

We strongly condemn any acts of corruption and violation of rules by way of offering or paying bribes or undue favours, either directly or indirectly. We adopt global best practices in corporate governance and business responsibility.

Besides our employees, all our consultants, suppliers and partners are also bound by a strong code of conduct that has a detailed clause on anti-bribery and anti-corruption.

All our product approvals are backed by science and clinical data.

The rationale for waiver of Phase 3 clinical trials was based on the following Indian regulatory guidance {Similar Biologics Guidelines 2016 & New Drugs and Clinical Trials 2019 (GSR 227 E)}.

The guidelines provide a framework for waiver of Phase 3 clinical trials to be conducted in India based on a commitment to undertake a Phase 4 trial, the *design of which* should be approved by the Central Licencing Authority.

In line with the above regulations, Biocon Biologics presented a proposal for import and marketing of Insulin Aspart with a waiver of Phase 3 clinical trial in India. The Company presented a detailed proposal along with CMC, pre-clinical and clinical trial data.

The Subject Expert Committee (Endocrinology and Metabolism) in its meeting held on May 18, 2022 at CDSCO, New Delhi, noted that Biocon Biologics has conducted Phase 1 and Phase 3 trials with Aspart in Germany and USA, respectively, and based on the



results of this global trial, Biocon Biologics' product, Aspart, has been granted marketing authorization by EMA and Health Canada.

"After detailed deliberation, the committee recommended for grant of permission to import and market the drug with waiver of Phase 3 clinical trial in the country with the condition that firm should conduct Phase 4 clinical trial in India (which also includes a sub-set population to generate PK/PD and immunogenicity and submit the protocol to CDSCO before placing the drug in the market) as per existing guidelines in the country," the SEC concluded.

[Source: Recommendations of the SEC (Endocrinology & Metabolism) made in its 87th meeting held on 18.05.2022 at CDSCO (HQ), New Delhi.

Link: <a href="https://cdsco.gov.in/opencms/opencms/system/modules/CDSCO.WEB/elements/common download.jsp?num id pk=MTY40Q==">https://cdsco.gov.in/opencms/opencms/system/modules/CDSCO.WEB/elements/common download.jsp?num id pk=MTY40Q==</a>]

Biocon Biologics follows due regulatory process for all our product approvals by the DCGI. The entire application process in India is online and all meeting minutes can be found on the website of the Central Drugs Standard Control Organization (CDSCO).

It is unfortunate that Biocon Biologics has been named in this controversy.

We reiterate that we strongly condemn all acts of bribery and corruption and have been co-operating with the investigation agency.

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