Subject: Company Statement

Dear Sir/Madam,

Please find below a Company Statement.

**Biocon Biologics’ New mAbs Facility Receives EU GMP Certification for bBevacizumab**

“Biocon Biologics’ integrated, multi-product, monoclonal antibodies (mAbs) Drug Substance manufacturing facility (B3) at Biocon Park, Bengaluru, has received a Certificate of GMP Compliance for an additional product, biosimilar Bevacizumab, from the representative European inspection authority, Health Products Regulatory Authority (HPRA), Ireland.

“This approval reflects Biocon Biologics’ compliance with the highest international regulatory standards and enables the Company to continue addressing the needs of patients in the EU through its high-quality products.

“This (B3) facility, which is one of India’s largest monoclonal antibodies (mAbs) manufacturing facilities, had received the EU GMP Certification for manufacturing biosimilar Trastuzumab last year. It was also awarded the ‘Facility of the Year Award’ (FOYA) with an ‘Honourable Mention’, by the International Society for Pharmaceutical Engineering (ISPE) in 2021”– Company Spokesperson

The above information will also be available on the website of the Company at [www.biocon.com](http://www.biocon.com).

Kindly take the same on record and acknowledge.

Thanking You,

Yours faithfully,

For Biocon Limited

Mayank Verma
Company Secretary and Compliance Officer
Membership No.: ACS 18776