

CE

## DECLARATION OF CONFORMITY

**Manufacturer:** Joinstar Biomedical Technology Co.,Ltd.

**Address:** 10th Floor, Administration Building, NO.519, XingGuo RD., Yuhang Economic and Technological Development Zone, Hangzhou, Zhejiang, China, 311188

**EC Representative's Name:** Lotus NL B.V.

**EC Representative's Address:** Koningin Julianaplein 10, 1e Verd, 2595AA, The Hague, Netherlands.

**Declares, that the product**

**Product Name and Model:**

COVID-19 Antigen Rapid Test (Latex)

1 Test/Kit, 5 Tests/Kit, 10 Tests/Kit, 20 Tests/Kit, 25 Tests/Kit.

**as described above are in conformity with the requirements as defined in IVDD98/79/EC Annex III.**

**Additional information:**


Conformity assessment route: Directive 98/79/EC, Annex III

Classification: List Others

I, the undersigned, hereby declare that the medical devices specified above conform with the Directive 98/79/EC on in vitro diagnostic medical devices and pertinent essential requirements.

Date Signed:

2021-09-16



Zhong WANG

Management Representative

Joinstar Biomedical Technology Co., Ltd.

Joinstar Biomedical Technology Co., Ltd.

