

EC DECLARATION OF CONFORMITY

Name and address of the manufacturer: **Hangzhou Clongene Biotech Co., Ltd.**
No.1 Yichuang Road, Yuhang Sub-district
Yuhang District
311121 Hangzhou
China

We declare under our sole responsibility that

the medical device: **COVID-19 Antigen Rapid Test**

of class: **Other**
according to article 9 of directive 98/79/EC

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the "final inspection report" of the device.

Conformity assessment procedure: **Directive 98/79/EC Annex III, excluding Section 6**

Applicable standards: **EN ISO 13485:2016** **EN ISO 15223-1:2016**
EN ISO 18113-1:2011 **EN ISO 18113-2:2011**
EN ISO 23640:2015 **EN ISO 14971:2019**
EN 13641:2002 **EN13612:2002/AC:2002**
EN 13975:2003 **EN 62366-1:2015**
ISO 17511:2020

Name and address of the authorized representative: **Shanghai International Holding Corporation GmbH (Europe)**
Eiffestrasse 80
20537 Hamburg
Germany



Hangzhou, February 22, 2021.

Place, date

Shujian Zheng, Legal representative

Name and function