

CE

CE

EC Declaration of Conformity

according to the Directive 98/79/EC

(applicable to IVD Devices of NOT Annex II and NOT self-test)

Manufacturer: Safecare Biotech (Hangzhou) Co., Ltd.

Address: Building 2/203, No.18 Haishu Rd.Cangqian Sub-district,
Yuhang District, Hangzhou, Zhejiang China 311121

EC Representative: NIC GmbH
Erlenweg 13,49076 Osnabrück, Germany

We, the manufacturer, declare under our sole responsibility that

the medical device(s) Product Name COVID-19 Antigen Rapid Test Kit(Swab)

Type/model, identification of product allowing traceability
(Where applicable) Cassette(COV Ag-6012)

of Category: Common/Others IVD
(Devices of NOT Annex II and NOT self-test)

is/are in conformity with the relevant provisions and requirements of Directive 98/79/EC of the European Parliament and of the Council on In-Vitro Diagnostic Medical Devices.

Applied harmonised standards, national standards or other normative documents	EN ISO23640:2015	EN ISO 18113-1:2011
	EN 13612:2002	ISO 18113-2: 2009
	EN 13641:2002	EN1041- 2008
	EN ISO 14971:2019	EN ISO15223-1:2016
	ISO13485:2016	

Conformity assessment procedure **Module A (EC Declaration of Conformity) (Annex III, except point 6)**

Notified Body (name & number) **NOT applicable**
Certificate & number

Signed on 28th Sep.,2020 Place: Hangzhou, Zhejiang, China

Signature (on behalf of the manufacturer) Kebin Qiu 2020.9.28

Name of authorized signatory: Kebin, Qiu
Position held in the company: General Manager

Seal/Stamp: 