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(Saliva)
	Type/model, identification of product allowing traceability (Where applicable)	COV Ag-7012
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- 2016
	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) Certificate & number	**NOT applicable**	

Signed on 3th Feb.,2021 Place: Hangzhou, Zhejiang, China

Signature (on behalf of the manufacturer) Kebin Qiu 2021.2.3

Name of authorized signatory: Kebin, Qiu

Position held in the company: General Manager

Seal/Stamp: