CE

DECLARATION OF CONFORMITY

According Directive 98/79/EC on in vitro diagnostic medical devices, Annex III.

Manufacturer: Hangzhou HEO Technology Co., Ltd

Address:(Room 201, Building 3, No. 2073 Jinchang Road, Yuhang District, Hangzhou,China

European Representative: Lotus NL B.V.

Contact person: Peter E-mail: peter@lotusnl.com

Address: Koningin Julianaplein 10,1e Verd, 2595AA,The Hague, Netherlands.

In Vitro Diagnostic Directive:

- COVID-19 Antigen Rapid Test Kit（Saliva）

Category: Others.

Conformity assessment route: Declaration of Conformity IVDD Annex III.

Applicable Standards:

ISO 13485:2016	*EN ISO 18113-3:2011*	*EN 13612:2002*
ISO 14971:2019	*EN 13641:2002*	*ISO 23640:2015*
EN ISO 18113-1:2011	*ISO 15223-1:2016*	*EN 62366-1:2015*
EN ISO 18113-2:2011		

We, the manufacturer, here declare with sole responsibility that our products mentioned above meets the provisions of the Directive 98/79/EC of the European Parliament and of the Council on In Vitro Diagnostic Medical Devices.

We agree to develop, implement and maintain a documented post-production monitoring process.

Signed: Tongwei, Sun

Place:Hangzhou,China

Name of authorized signatory: Tongwei Sun

Position held in the company: General Manager

Date: Aug-31th, 2021

Seal/Stamp:

Hangzhou HEO Technology Co., Ltd.