DECLARATION OF CONFORMITY

According to the *In vitro* Diagnostic Medical Device Directive 98/79/EC

Manufacturer	Guangdong Longsee Biomedical Co.,Ltd.
Address	5/F Building A, No.83, Ruihe Road, Huangpu District, 510000, Guangzhou, China
European Representative	Qarad EC-REP BV
Address	Pas 257, 2440 Geel, Belgium
Product Information	2019-nCoV Ag Rapid Detection Kit (Immuno-Chromatography)
Type	LS-C-T-008(U3), LS-C-T-008(V3)
Classification	IVD for self-testing
Conformity Assessment Route	IVDD Annex IV, excluding 4 and 6

General Applicable Directives: *In vitro* Diagnostic Medical Device Directive 98/79/EC

Standards Applied	EN 13612:2002	EN 13612:2002/AC:2002
	EN ISO 23640:2015	EN 13641:2002
	EN ISO 14971:2012	EN ISO 15223-1:2016
	EN ISO 18113-1:2011	EN ISO 18113-4:2011
	EN ISO 13485:2016	EN ISO 13485:2016/AC:2018
	EN 13532:2002	EN 62366-1:2015

We,the manufacturer, hereby declare under our sole responsibility that the above mentioned products meet the provisions of the above EC Council Directives and Standards. The products meet prospective uses and all supporting documentations are retained under the premises of the manufacturer.

Notified Body	Polish Centre for Testing and Certification (NB 1434)
Address	469 Putawska Street, 02-844 Warsaw, Poland
EC Certificate No.	1434-IVDD-014/2022
Expiry date of the Certificate	May 27, 2025

Place, date of issued: Guangzhou, P. R. China, May 1, 2022

Signature of Chief Executive Officer:

Shizhou Deng

(Shizhou Deng)

CE 1434