CERTIFICATE

EC Certificate No. 1434-IVDD-014/2022

EC Design-examination
Directive 98/79/EC concerning
***in vitro* diagnostic medical devices**

Polish Centre for Testing and Certification certifies
that manufactured by:

Guangdong Longsee Biomedical Co.,Ltd.
5/F Building A, No.83, Ruihe Road, Huangpu District, 510000, Guangzhou, China

in vitro diagnostic medical devices
for self-testing

2019-nCoV Ag Rapid Detection Kit (Immuno-Chromatography)
Ref. No.: LS-C-T-008 (U3), LS-C-T-008 (V3)

in terms of design documentation, comply with requirements
of Annex III (Section 6) to Directive 98/79/EC (as amended)
implemented into Polish law,
as evidenced by the audit conducted by the PCBC

Validity of the Certificate: from 01.02.2022 to 27.05.2025

The date of issue of the Certificate: 01.02.2022

The date of the first issue of the Certificate: 01.02.2022

CE 1434

Issued under the Contract No. MD-68/2021
Application No: 150/2021
Certificate bears the qualified signature.
Warsaw, 01/02/2022
Module A1

Aleksandra Kostrzewa
Digitally signed by Aleksandra Kostrzewa
President