CERTIFICATE

EC Certificate No. 1434-IVDD-216/2022

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

Polish Centre for Testing and Certification certifies
that manufactured by:

Hangzhou AllTest Biotech Co., Ltd.

550#, Yinhai Street, Hangzhou Economic and Technological Development Area, Hangzhou- 310018, P.R. China

in vitro diagnostic medical devices
for self-testing

SARS-CoV-2 (COVID-19) Antigen Rapid Test (Saliva)

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 24.05.2022 to 27.05.2025

The date of issue of the Certificate: 24.05.2022

The date of the first issue of the Certificate: 24.05.2022

CE 1434

Issued under the Contract No. MD-162/2021
Application No: 309/2021
Certificate bears the qualified signature.
Warsaw, 24/05/2022
Module A1

Tomasz Koeber
Elektronicznie podpisany przez Tomasz Artur Koeber
Data: 2022.05.24 11:19:29 +02'00'

Director
Medical Devices Certification Department