CERTIFICATE

EC Certificate No. 1434-IVDD-193/2022

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

Polish Centre for Testing and Certification certifies
that manufactured by:

Jiangsu Medomics Medical Technology Co., Ltd.
F3, Building C, No.3-1 Xinjinhu Road,
Jiangbei New Area, Nanjing, Jiangsu 210030 CHINA

in vitro diagnostic medical devices
for self-testing

SARS-CoV-2 & Influenza A/B Antigen Combo Rapid Test Kit (LFIA)
Ref. no.: 1041-14-01, 1041-24-01, 1041-34-01, 1041-54-01

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-206/2021
Application No: 582/2021
Certificate bears the qualified signature.
Warsaw, 24/05/2022
Module A1

Tomasz Koeber

Elektronicznie podpisany
przez Tomasz Artur Koeber
Data: 2022.05.24 07:10:38
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Director
Medical Device Certification
Department