

EC Certificate No. 1434-IVDD-217/2022

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 and Influenza A+B Antigen Combo Rapid Test (Nasal Swab)

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



Issued under the Contract No. MD-232/2021 Application No: 465/2021 Certificate bears the qualified signature. Warsaw, 24/05/2022 Module A1 Tomic Koeta

Elektronicznie podpisany przez Tomasz Artur Koeber Ostar 2022.05.24 11:19:56 +02:00*

Director Medical Devices Certification Department