CERTIFICATE

EC Certificate No. 1434-IVDD-068/2022

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

Polish Centre for Testing and Certification certifies
that manufactured by:

Anbio (Xiamen) Biotechnology Co., Ltd.
No.2016, Wengjiao West Road, Xinyang Street, Haicang District, 361026 Xiamen, Fujian, China

in vitro diagnostic medical devices
for self-testing

Rapid COVID-19 Antigen Test (Colloidal Gold) / Nasal Swab

A606101, A606102, A606103, A606104, A606105, A606106

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 30.03.2022 to 27.05.2025

The date of issue of the Certificate: 30.03.2022

The date of the first issue of the Certificate: 16.08.2021

CE 1434

Issued under the Contract No. MD-124/2021
Application No: 243/2021
Certificate bears the qualified signature.
Warsaw, 30/03/2022
Module A1

Aleksandra Kostrzewa
Digitally signed by Aleksandra Kostrzewa
President