CERTIFICATE

EC Certificate No. 1434-IVDD-467/2021

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

Polish Centre for Testing and Certification certifies
that manufactured by:

Xiamen AmonMed Biotechnology Co., Ltd

Unit 503, 120 Xinyuan Road, Haicang District,
Xiamen, Fujian, China

in vitro diagnostic medical devices
for self-testing

COVID-19 Antigen Rapid Test Kit (Colloidal Gold)
Saliva specimen

CG01Ag-01S-ST, CG01Ag-05S-ST, CG01Ag-25S-ST

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 15.10.2021 to 27.05.2024

The date of issue of the Certificate: 15.10.2021

The date of the first issue of the Certificate: 15.10.2021

CE 1434

Issued under the Contract No. MD-128/2021
Application No: 233/2021
Certificate bears the qualified signature.
Warsaw, 15/10/2021
Module A1
FBM-30-E_10

Anna Małgorzata Wyroba

Elektronicznie podpisany przez Anna Małgorzata Wyroba
Data: 2021.10.15 13:27:16 +02'00'

Vice-President