CERTIFICATE

EC Certificate No. 1434-IVDD-157/2022

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

Polish Centre for Testing and Certification certifies
that manufactured by:

Shenzhen Lvshiyuan Biotechnology Co., Ltd

101, 201, 301 D Building, No. 2 Industrial Avenue, Buxin Village, Buxin Community, Dapeng Subdistrict Office, Dapeng New District, Shenzhen 518120, China

in vitro diagnostic medical devices
for self-testing

SARS-CoV-2 Antigen Rapid Test Kit (Colloidal Gold)

Ref. No.: GF102BS1, GF102BS5, GF102BS10, GF102BS25

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 16.05.2022 to 27.05.2025

The date of issue of the Certificate: 16.05.2022

The date of the first issue of the Certificate: 16.05.2022

CE 1434

Issued under the Contract No. MD-43/2021
Application No: 94/2021
Certificate bears the qualified signature.
Warsaw, 16/05/2022
Module A1

Aleksandra Kostrzewa
Digitally signed by Aleksandra Kostrzewa
President