CERTIFICATE

EC Certificate No. 1434-IVDD-149/2022

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

Polish Centre for Testing and Certification certifies
that manufactured by:

Anhui Deepblue Medical Technology Co., Ltd. 4th Floor, D-1#Zone, Pearl Industrial Park, 106 Innovation Avenue, High-Tech Development Zone, 230088 Hefei, Anhui, People's Republic Of China

in vitro diagnostic medical devices
self-testing

COVID-19 (SARS-CoV-2) Antigen Test Midstream – Saliva

The list of medical devices covered by this certificate is provided in the annex 1

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 11.05.2022 to 27.05.2025

The date of issue of the Certificate: 11.05.2022

The date of the first issue of the Certificate: 11.05.2022

CE 1434

Issued under the Contract No. MD-159/2021
Application No: 291/2021
Certificate bears the qualified signature.
Warsaw, 11/05/2022
Module A1

President

ANNEX 1 TO THE CERTIFICATE

VALID ONLY WITH CERTIFICATE

No 1434-IVDD-149/2022

List of medical devices covered by the certificate:

Brands names	*Ref No.*
DEEPBLUE, ShenLanTest, Highbluetest, NewBluetest, Quicklyblue the one medical in.pro.Medical	COVAg3SST-1
	COVAg3SST-2
	COVAg3SST-3
	COVAg3SST-5
	COVAg3SST-6
	COVAg3SST-7
	COVAg3SST-8
	COVAg3SST-9
	COVAg3SST-10
	COVAg3SST-11
	COVAg3SST-12
	COVAg3SST-15
	COVAg3SST-16
	COVAg3SST-17
	COVAg3SST-18
	COVAg3SST-19
	COVAg3SST-20
	COVAg3SST-25

CE 1434

Issued under the Contract No. **MD-159/2021**
Application No: **291/2021**
Certificate bears the qualified signature.
Warsaw, 23/05/2022

Director
Medical Device
Certification
Department