CERTIFICATE

EC Certificate No. 1434-IVDD-261/2022

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

Polish Centre for Testing and Certification certifies
that manufactured by:

Hangzhou GENESIS Biodetection and Biocontrol Co., Ltd.

No. 139, 10 th Street(East), Hangzhou Economic&Technological Development Zone, 310018 Hangzhou, China

in vitro diagnostic medical devices
for self-testing

EZER Flu & COVID-19 Antigen Duo Rapid Test

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 25.05.2022 to 27.05.2025

The date of issue of the Certificate: 25.05.2022

The date of the first issue of the Certificate: 25.05.2022

CE 1434

Issued under the Contract No. MD-231/2021
Application No: 442/2021
Certificate bears the qualified signature.
Warsaw, 25/05/2022
Module A1

Tomasz Koeber

Elektronicznie podpisany
przez Tomasz Artur Koeber
Data: 2022.05.25 22:27:41
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Director
Medical Devices Certification Department