CERTIFICATE

EC Certificate No. 1434-IVDD-031/2022

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

Polish Centre for Testing and Certification certifies
that manufactured by:

Rapid Labs Ltd
Unit 2 & 2a Hall Farm Business Centre; Church Road; Little Bentley; Colchester Essex; CO7 8SD; United Kingdom

i.e. *in vitro* diagnostic medical devices
List A

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 IV (Section 4) 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 16.03.2022 to 27.05.2025
The date of issue of the Certificate: 16.03.2022

The date of the first issue of the Certificate: 08.03.2022

CE 1434

Issued under the Contract No. MD-29/2021
Application No: 515/2021
Certificate bears the qualified signature.
Warsaw,16/03/2022

Aleksandra Kostrzewa
President

Digitally signed by Aleksandra Kostrzewa

ANNEX 1 TO THE CERTIFICATE

VALID ONLY WITH CERTIFICATE

No 1434-IVDD-031/2022

List of medical devices covered by the certificate:

Anti-A,B Monoclonal 10ml BG-AB10

Anti-A,B Monoclonal 10 x 10ml BG-AB10X10

Anti-A Monoclonal 10ml BG-A10

Anti-A Monoclonal 10 x 10ml BG-A10X10

Anti-B Monoclonal 10ml BG-B10

Anti-B Monoclonal 10 x 10ml BG-B10X10

Anti-D (IgG & IgM) Monoclonal 10ml BG-D10

Anti-D (IgG & IgM) Monoclonal 10 x 10ml BG-D10X10

CE 1434

Issued under the Contract No. MD-29/2021
Application No: 515/2021
Certificate bears the qualified signature.
Warsaw, 16/03/2022

Aleksandra Kostrzewa
President

Digitally signed by Aleksandra Kostrzewa