

EC Certificate No. 1434-IVDD-001/2022

Full Quality Assurance System
Directive 98/79/EC concerning
in vitro diagnostic medical devices

Polish Centre for Testing and Certification certifies that the quality assurance system in the organization:

ProBioQual

7 Rue Antoine Lumiére 69008 Lyon, France

for the design, manufacture and final inspection of *in vitro* diagnostic medical device List B

CIQ - MSMPE - Sérum S

complies with requirements of Annex IV (excluding Section 4, 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 01.01.2022 to 27.05.2025

The date of issue of the Certificate: 29.12.2021

The date of the first issue of the Certificate: 23.06.2020



Issued under the Contract No. MD-172/2021 Application No: 571/2021 Certificate bears the qualified signature. Warsaw, 29/12/2021 Module H7

FBM-32-E_9

President