

EC Certificate No. 1434-IVDD-404/2020 EC Design-examination

Directive 98/79/EC concerning in vitro diagnostic medical devices

Polish Centre for Testing and Certification certifies that manufactured by:

TÜRKLAB Tıbbi Mal. San. Tic. A.S. ITOB 10031 Sokak No: 15, Tekeli - Menderes Izmir, Turkey

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

Blood Grouping Gel Cards Reagent Red Blood Cells

The list of medical devices covered by this certificate is provided in the annex 1 to EC Design-examination Certificate No. 1434-IVDD-403/2020

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 03.11.2020 to 27.05.2024

The date of issue of the Certificate: 03.11.2020



Issued under the Contract No. MD-10/2020 Application No: 447/2019, 449/2019 Certificate bears the qualified signature. Warsaw,03.11.2020 Module H6

Vice-President