# CERTIFICATE

## EC Certificate No. 1434-IVDD-252/2022

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

Polish Centre for Testing and Certification certifies
that manufactured by:

## Vitrosens Biyoteknoloji LTD. Şti
## Serifali Mah. Sehit Sok. No: 17 34775 Istanbul, Turkey.

in vitro diagnostic medical devices
for self-testing

## RapidFor™ SARS-CoV-2 Rapid Antigen Test Kit (Nasal)
## 1Test/Box – REF. VSCD02ST01
## 2 Tests/Box – REF. VSCD02ST02
## 5 Tests/Box – REF. VSCD02ST05

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-203/2021
Application No: 422/2021
Certificate bears the qualified signature.
Warsaw, 25/05/2022
Module A1

Tomasz Koeber
Elektronicznie podpisany
przez Tomasz Artur Koeber
Data: 2022.05.25 12:33:39
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**Director**
**Medical Device Certification**
**Department**