

EC Certificate No. 1434-IVDD-054/2022

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

Polish Centre for Testing and Certification certifies that manufactured by:

Citest Diagnostics Inc. 170-422 Richards Street, Vancouver, CANADA

in vitro diagnostic medical devices for self-testing

COVID-19 Antigen Rapid Test (Swab) REF. No. ICOV-502S

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 23.03.2022 to 27.05.2025

The date of issue of the Certificate: 23.03.2022

The date of the first issue of the Certificate: 25.11.2021



Issued under the Contract No. MD-51/2021 Application No: 140/2021 Certificate bears the qualified signature. Warsaw, 23/03/2022 Module A1

President