

EC Certificate No. 1434-IVDD-483/2021

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

Polish Centre for Testing and Certification certifies that manufactured by:

Goldsite Diagnostics Inc.
No. 103C, 503C & 504D, Technology Building & No. 3A & 4A, Technology Building Annex, Zhaoshang Sub-District,
Nanshan District,
Shenzhen, China

in vitro diagnostic medical devices self-testing

SARS-CoV-2 Antigen Kit (Colloidal Gold)

REF: CG123001, CG123003, CG123005, CG123007, CG123025

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 10.11.2021 to 27.05.2024

The date of issue of the Certificate: 10.11.2021

The date of the first issue of the Certificate: 10.11.2021



Issued under the Contract No. MD-40/2021 Application No: 397/2020 Certificate bears the qualified signature. Warsaw, 10/11/2021 Module A1

Vice-President *Mgr. Anna Wyroba*