



CERTIFICATE

EC Certificate No. 1434-IVDD-492/2021

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

Polish Centre for Testing and Certification certifies
that manufactured by:

**Xiamen Wiz Biotech Co., Ltd.
3-4 Floor, NO.16 Building, Bio-medical Workshop,
2030 Wengjiao Xi Road, Haicang District, Xiamen City,
Fujian Province, 361026, P.R. China**

in vitro diagnostic medical devices
for self-testing

SARS-CoV-2 Antigen Rapid Test (Colloidal Gold)

51332801, 51332802, 51332803, 51332804

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 22.11.2021 to 27.05.2024

The date of issue of the Certificate: 22.11.2021

The date of the first issue of the Certificate: 22.11.2021



Issued under the Contract No. **MD-77/2021**
Application No: **130/2021**
Certificate bears the qualified signature.
Warsaw, 22/11/2021
Module A1

Vice-President