

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 Anna Małgorzata Wyroba

Elektronicznie podpisany przez Anna Małgorzata Wyroba Data: 2021.11.22 13:16:54 +01'00'

Vice-President