

EC Certificate No. 1434-IVDD-491/2021

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

Polish Centre for Testing and Certification certifies that manufactured by:

Shenzhen Microprofit Biotech Co., Ltd., Rm. 405, 406, Zone B /4F, Rm. 205, 206-1, 207, West Side of Zone B/ 2F, Haowei Building, No. 8 Langshan 2nd Road, Songpingshan, Songpingshan Community, Xili Street, Nanshan District, Shenzhen, P.R. China

in vitro diagnostic medical devices for self-testing

SARS-CoV-2 Antigen Test Kit (Colloidal Gold Chromatographic Immunoassay) REF: MF-68

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-76/2021 Application No: 111/2021 Certificate bears the qualified signature. Warsaw, 22/11/2021 Module A1

Vice-President