



CERTIFICATE

EC Certificate No. 1434-IVDD-454/2021

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

Polish Centre for Testing and Certification certifies
that manufactured by:

**Anbio (Xiamen) Biotechnology Co., Ltd.
No.2016, Wengjiao West Road, Xinyang Street, Haicang
District, 361026 Xiamen, Fujian, China.**

in vitro diagnostic medical devices
for self-testing

**Rapid COVID-19 Antigen Test (Colloidal Gold) / Saliva
A606201/A606202/A606203/A606204/A606205/606206**

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 31.08.2021 to 27.05.2024

The date of issue of the Certificate: 31.08.2021

The date of the first issue of the Certificate: 31.08.2021



Issued under the Contract No. MD-124/2021
Application No: 243/2021
Certificate bears the qualified signature.
Warsaw, 31/08/2021
Module A1

Vice-President
Mgr Anna Wyroba