CERTIFICATE

DIRECTIVE 98/79/EC
EC DESIGN-EXAMINATION

CeCert Sp. z o.o. hereby confirms that manufactured by

Ningbo Lvtang Biotechnology Co., Ltd.
No. 9 Dongpu Road, Chengdong Industrial Area,
Daxu Town, Xiangshan County, Ningbo City,
Zhejiang Province, P.R. China

in vitro diagnostic medical device for self-testing

SARS-CoV-2 Antigen Detection Kit (Colloidal Gold Method)
catalogue numbers: CovAg-N-1, CovAg-N-2, CovAg-N-5, CovAg-N-20, CovAg-N-25

in term of the design conforms to the requirements of Annex III section 6 to Directive 98/79/EC (as amended) implemented into Polish Law, as evidenced by the assessment conducted by CeCert Sp. z o.o.

CE 2934

Validity date: 29.04.2022 – 26.05.2025

Issue date: 29.04.2022

Check it

Kamil Szczurowski
Director of *in Vitro* Diagnostic Medical Device
Certification Department

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Certificate no: CeCert/061/W/E.1