CERTIFICATE

DIRECTIVE 98/79/EC
EC DESIGN-EXAMINATION

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

Hangzhou Lysun Biotechnology Co., Ltd.
6th Floor, 6th Building, No. 95 Binwen Road, Xixing Street, Binjiang District, Hangzhou, Zhejiang, P.R. China

in vitro diagnostic medical device for self-testing

COVID-19 Antigen Rapid Test Device (Colloidal Gold)
catalogue numbers: 3040102201, 3040102301, 3040102101

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: 17.05.2022 – 26.05.2025

Issue date: 17.05.2022

Check it

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

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Certificate no: CeCert/105/W/E.1