

CERTIFICATE

DIRECTIVE 98/79/EC EC DESIGN-EXAMINATION

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

Suny (Xiamen) Biotechnology Co., Ltd.

6th Floor, Unit 3 and 7th Floor, Unit 3, Building B1, No. 2050, Wengjiao West Road, Haicang District, 361026 Xiamen, Fujian, P.R. China

in vitro diagnostic medical device for self-testing

Rapid SARS-CoV-2 & Flu A & Flu B Antigen Test (Colloidal Gold)

catalogue numbers: \$6061701, \$6061702, \$6061703, \$6061704

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.

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Validity date: 06.05.2022 - 26.05.2025

Issue date: 06.05.2022

Check it

CeCert Sp. z o.o. ul. Żurawia 32/34 00-515 Warszawa MUUN VALU Kamil Szczurowski

Director of in Vitro Diagnostic Medical Device Certification Department