CERTIFICATE

DIRECTIVE 98/79/EC
EC DESIGN-EXAMINATION

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

Hangzhou Realy Tech Co., Ltd.
#2 Building, No. 763, Yuansha Village
Xinjie Street, Xiaoshan District
311200 Hangzhou City, Zhejiang Province, P.R. China

in vitro diagnostic medical device for self-testing

Novel Coronavirus (SARS-Cov-2) Antigen Rapid Test Device (nasal swab), catalogue number: K601416D

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: 02.03.2022 – 26.05.2025

Issue date: 02.03.2022

Check it

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

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Certificate no: CeCert/025/W/E.1