CERTIFICATE

DIRECTIVE 98/79/EC
EC DESIGN-EXAMINATION

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

Koch Biotechnology (Beijing) Co., Ltd
No. 16, Chunlin Street, Daxing District,
Beijing, P.R. China

in vitro diagnostic medical device for self-testing

COVID-19 Antigen Rapid Test Kit
catalogue numbers:
NCV12-SEU, NCV12-SEUA, NCV13-SEU, NCV11-SEU

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: 09.05.2022 – 26.05.2025

Issue date: 09.05.2022

Check it

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

CeCert Sp. z o.o.
ul. Żurawia 32/34
00-515 Warszawa

www.cecert.pl
e-mail: biuro@cecert.pl

Certificate no: CeCert/080/W/E.1