CeCert.

CERTIFICATE

DIRECTIVE 98/79/EC
EC DESIGN-EXAMINATION

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

Biobase Diagnostic Technology (Shandong) Co., Ltd.
A402, Floor 4, Building #2, BIOBASE Headquarters,
No. 9 Gangxing Road, Pilot Free Trade Zone of Jinan,
Shandong, P.R. China

in vitro diagnostic medical device for self-testing

Biobase COVID-19 Antigen Rapid Test
catalogue numbers:
DR001-1, DR001-2, DR001-3, DR001-4

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: 15.04.2022 – 26.05.2025

Issue date: 15.04.2022

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/047/W/E.1