CERTIFICATE

DIRECTIVE 98/79/EC
EC DESIGN-EXAMINATION

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

Xiamen Hopegen Medical Technology Co., Ltd.

Room 905, 253 Duiying Nan Road, Houxi Town,
Jimei District, Xiamen, Fujian, P.R. China

in vitro diagnostic medical device for self-testing

COVID-19 Antigen Rapid Test Kit (Colloidal Gold)

catalogue numbers:
XJ-ZC-411, XJ-ZC-412, XJ-ZC-413, XJ-ZC-414

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: 24.05.2022 – 26.05.2025

Issue date: 24.05.2022

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/127/W/E.1