CERTIFICATE

DIRECTIVE 98/79/EC
EC DESIGN-EXAMINATION

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

LOMINA Superbio a.s.
Na Radosti 184/59, Prague 5,
155 21, Czech Republic

in vitro diagnostic medical device for self-testing

Lomina SARS-CoV-2
Antigen LTX Selftest
catalogue numbers:
L-LTX-ST/1P, L-LTX-ST/1B, L-LTX-ST/2B,
L-LTX-ST/25B, L-LTX-ST/50B

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: 10.05.2022 – 26.05.2025

Edition issue date: 26.07.2022

Check it

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

CeCert Sp. z o.o.
ul. Warecka 11A
00-034 Warszawa

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

Certificate no: CeCert/083/W/E.2