

## **CERTIFICATE**

DIRECTIVE 98/79/EC
EC DESIGN-EXAMINATION

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

Safecare Biotech (Hangzhou) Co., Ltd.

Building 2/203, No. 18 Haishu Rd., Cangqian Sub-district, Yuhang District, Hangzhou, 311121, Zhejiang, P.R. China

in vitro diagnostic medical device for self-testing

COVID-19 & Influenza A+B Antigen Combo Rapid Test

catalogue number: FCO-6032H

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.

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Validity date: 29.04.2022 - 26.05.2025 Issue date: 29.04.2022

Check it

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Kamil Szczurowski Director of in Vitro Diagnostic Medical Device Certification Department

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Certificate no: CeCert/063/W/E.1



