CERTIFICATE

DIRECTIVE 98/79/EC
EC DESIGN-EXAMINATION

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

Shenzhen Microprofit Biotech Co., Ltd.
Rm. 405, 406, Zone B/4F, Rm. 205, 206-1, 207, West Side of Zone B/ 2F, Haowei Building, No. 8 Langshan 2nd Road, Songpingshan, Songpingshan Community, Xili Street, Nanshan District, Shenzhen, P.R. China

in vitro diagnostic medical device for self-testing

fluorecare SARS-CoV-2 & Influenza A/B & RSV Antigen Combo Test Kit
catalogue numbers: MF-71-1, MF-71-2, MF-71-5

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: 12.05.2022 – 26.05.2025

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Check it

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

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Certificate no: CeCert/092/W/E.2