# CERTIFICATE

## DIRECTIVE 98/79/EC
## EC DESIGN-EXAMINATION

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

**InTec PRODUCTS, INC.**
**332 Xinguang Road, Xinyang Industrial Area,**
**Haicang, 361022, Xiamen, Fujian, P.R. China**

*in vitro* diagnostic medical device for self-testing

**Rapid SARS-CoV-2 Antigen Test**
**catalogue numbers:**
**ITP16030-TC1, ITP16030-TC2, ITP16030-TC5,**
**ITP16030-TC7, ITP16030-TC25**

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: 04.04.2022 – 26.05.2025

Issue date: 04.04.2022

Check it

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

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Certificate no: CeCert/034/W/E.1