Digitally signed by Jighili Tatiana
Date: 2024.11.08 17:04:59 EET
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MOLDOVA EUROPEANĂ

CERTIFICATE

DIRECTIVE 98/79/EC
EC DESIGN-EXAMINATION

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

Hangzhou AllTest Biotech Co., Ltd.
#550, Yinhai Street, Hangzhou Economic & Technological Development Area, Hangzhou, 310018, P.R. China

in vitro diagnostic medical device referred to in List A in Annex II

HCV Rapid Test Cassette (Whole Blood/Serum/Plasma)
catalogue number: IHC-402

in term of the design conforms to the requirements of Annex IV section 4 to Directive 98/79/EC (as amended) implemented into Polish Law, as evidenced by the audit conducted by CeCert Sp. z o.o.

CE 2934

Validity date: 17.05.2022 – 26.05.2025

Issue date: 17.05.2022

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/106/W/E.1

CERTIFICATE

DIRECTIVE 98/79/EC
FULL QUALITY ASSURANCE SYSTEM

CeCert Sp. z o.o. hereby confirms that the quality assurance system in the organization

Hangzhou AllTest Biotech Co., Ltd.
#550, Yinhai Street, Hangzhou Economic & Technological Development Area, Hangzhou, 310018, P.R. China

with regard to the design, manufacture and final inspection of *in vitro* diagnostic medical device referred to in List A in Annex II

HCV Rapid Test Cassette (Whole Blood/Serum/Plasma)
catalogue number: IHC-402

conforms to the requirements of Annex IV (excluding section 4 and 6) to Directive 98/79/EC (as amended) implemented into Polish Law, as evidenced by the audit conducted by CeCert Sp. z o.o.

CE 2934

Validity date: 17.05.2022 – 26.05.2025

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Certificate no: CeCert/107/W/E.1