Certificate

Certificate No.:	MD 3333169-1011
Manufacturer:	**Immucor GTI Diagnostics, Inc.** 20925 Crossroads Circle Waukesha WI 53186 USA
REPs Facility ID:	F003971
Certification criteria:	ISO 13485:2016 Australia Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure Brazil RDC ANVISA n. 16/2013, RDC ANVISA n. 23/2012, RDC ANVISA n. 67/2009 Canada Medical Devices Regulations – Part 1 – SOR 98/282 United States 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 – Subparts A to D
Scope:	Design, development and manufacture of in vitro diagnostic medical devices, reagents and software used in the management of immune status, coagulation, tissue and immunological typing.

Digitally signed by Talpiş Andrei
Date: 2023.09.24 21:10:33 EEST
Reason: MoldSign Signature
Location: Moldova

TUV Rheinland of North America, Inc., an MDSAP recognized Auditing Organization, certifies that the quality management system of the Manufacturer has been audited against and found to conform the Certification criteria for the Scope contained in this certificate. The quality management system is subject to annual surveillance audit(s).

Project No.:	3333169-160
Issue Date:	2021-12-13
Effective Date:	2022-01-04
Expiry Date:	2024-01-03

MDSAP
MEDICAL DEVICE SINGLE AUDIT PROGRAM

Certification officer: Dipl.-Ing. (FH) D. Wiedemuth
TUV Rheinland of North America, Inc.

The validity of the certificate can be verified on https://www.certipedia.com/quality_marks/9000009904?locale=en or calling 1-888-743-4652.

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