EU Certificate

Quality Management System
REGULATION (EU) 2017/746 on In Vitro Diagnostic Medical Devices, Annex IX Chapter I, Section 2 and 3 and Chapter III

TÜVRheinland®

Registration No.: HX 1804147-1

Manufacturer:
Immucor, Inc.
3130 Gateway Drive
Norcross GA 30071
USA

EUDAMED Single Registration No.: US-MF-000011568

Products: Products of class C

HAEMATOLOGY / HAEMOSTASIS / IMMUNOHAEMATOLOGY / HISTOLOGY / CYTOLOGY

IVR 106: Other devices intended to be used for blood grouping
W01030303 - ANTIBODY DETECTION (IMMUNOHAEMATOLOGY)
W01030304 - IMMUNOHAEMATOLOGY CONTROLS
W01030305 - OTHER ANTIGEN TYPING REAGENTS
W01030399 - IMMUNOHAEMATOLOGY (BLOOD GROUPING) TESTS - OTHER

IVR 201: Devices intended to be used for tissue typing (HLA A, B, DR) to ensure the immunological compatibility of blood, blood components, cells, tissue or organs that are intended for transfusion or transplantation or cell administration
W01030399 - IMMUNOHAEMATOLOGY (BLOOD GROUPING) TESTS – OTHER

The Notified Body hereby declares that the requirements of Annex IX, Chapter I, Section 2 and 3 of the REGULATION (EU) 2017/746 have been met for the listed products. The above named manufacturer has established and applies a quality management system, which is subject to periodic surveillance, defined by Annex IX, Chapter I, Section 3 of the aforementioned regulation. If class B, C or D devices for self-testing or near-patient testing are covered by this certificate an EU technical documentation assessment certificate according to Chapter II, Section 5.1 is required before placing them on the market. If companion diagnostics are covered by this certificate an EU technical documentation assessment certificate according to Chapter II, Section 5.2 is required before placing them on the market. If class D devices are covered by this certificate an EU technical documentation assessment certificate according to Chapter II, Section 4.10 is required before placing them on the market.

Report No.: 1132944-60

Effective date: 2023-09-19

Expiry date: 2028-05-10

Issue date: 2023-09-19

TÜV Rheinland LGA Products GmbH TÜVRheinland® Zertifizierstelle

Benannt durch/Designated by Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten www.zlg.de
BS-IVDR-097

Katja Mierisch
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

TÜV Rheinland LGA Products GmbH is a Notified Body according to REGULATION (EU) 2017/746 concerning in vitro diagnostic medical devices with the identification number 0197.

EU Certificate

Quality Management System
REGULATION (EU) 2017/746 on In Vitro Diagnostic Medical Devices, Annex IX Chapter I, Section 2 and 3 and Chapter III

TÜVRheinland®

Registration No.: HX 1804147-1

Manufacturer:

Immucor, Inc.
3130 Gateway Drive
Norcross GA 30071
USA

Products of class D

HAEMATOLOGY / HAEMOSTASIS / IMMUNOHAEMATOLOGY / HISTOLOGY / CYTOLOGY

IVR 0102: Devices intended to determine markers of the Rhesus system [RH1 (D), RHW1, RH2 (C), RH3 (E), RH4 (c), RH5 (e)]
W01030304 - IMMUNOHAEMATOLOGY CONTROLS

IVR 0104: Devices intended to determine markers of the Kidd system [JK1 (Jka), JK2 (Jkb)]
W01030304 - IMMUNOHAEMATOLOGY CONTROLS

IVR 0105: Devices intended to determine markers of the Duffy system [FY1 (Fya), FY2 (Fyb)]
W01030304 - IMMUNOHAEMATOLOGY CONTROLS

IVR 0106: Other devices intended to be used for blood grouping
W01030304 - IMMUNOHAEMATOLOGY CONTROLS

Authorised representative(s):

Immucor Medizinische Diagnostik GmbH
Robert-Bosch-Strasse 32
63303 Dreieich
Germany

Certificate history		
Revision:	Description:	Issue date:
0	Initial certification	2023-05-11
1	Scope extension, class D <<Weak D Cells>>	2023-09-19

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