

EC Certificate

**Full Quality Assurance System
Directive 98/79/EC on In Vitro Diagnostic Medical Devices,
Annex IV excluding (4, 6)**

Registration No.: HL 1804147-1

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

Products: Annex II List A Products:

- corQC Test System
- corQC EXTEND Standard
- corQC EXTEND 1, 2 and 3
- corQC EXTEND Complete
- Monoclonal Control
- Weak D cells
- Referencells-4 (Group A1, A2, B and O)
- Referencells-2 (Group A1 and B)
- Referencells-1 (Group A2)
- WB corQC

Annex II List B Products:

- Checkcell
- Checkcell (Weak)
- Panoscreen I and II
- Panoscreen I, II and III

The Notified Body hereby declares that the requirements of Annex IV, excluding section 4 and 6 of the directive 98/79/EC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex IV, section 5 of the aforementioned directive. For placing on the market of List A devices covered by this certificate an EC design-examination certificate according to Annex IV, section 4 and a verification of manufactured products according to section 6 is required.

Report No.: 1111389-10
Effective date: 2020-12-17
Expiry date: 2025-05-26
Issue date: 2022-05-10



Wenxiang Zhang
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 98/79/EC concerning in vitro diagnostic medical devices with the identification number 0197.

EC Certificate



Full Quality Assurance System
Directive 98/79/EC on In Vitro Diagnostic Medical Devices,
Annex IV excluding (4, 6)

Registration No.: HL 1804147-1

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

Panoscreen EXTEND
Panocell-16
Panocell-20
Panocell-10
Panocell-10, Ficin-Treated
Hemantigen
Capture-CMV (Plates)
Capture-CMV Indicator Red Cells
Capture-R Ready-Screen (I and II)
Capture-R Ready-Screen (Pooled Cells)
Capture-R Ready-ID
pHix

Capture-R Ready-ID EXTEND I
Capture-R Ready-ID EXTEND II
Capture-R Ready-Screen (3)
Anti-Jka (Monoclonal) Gamma-clone®
Anti-Jkb (Monoclonal) Gamma-clone®
Gamma PeG
Capture-R Ready-Screen (3)
Capture-R Ready-Screen (4)

Anti-Human Globulin, Anti-IgG, -C3d
Polyspecific (Murine Monoclonal) Gamma-Clone®
Anti-Human Globulin, Anti-IgG (Murine Monoclonal)
Gamma-Clone®

Report No.: 1111389-10

Effective date: 2020-12-17

Expiry date: 2025-05-26

Issue date: 2022-05-10



A handwritten signature in blue ink, appearing to read 'Zhang'.

Wenxiang Zhang
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 98/79/EC concerning in vitro diagnostic medical devices with the identification number 0197.

EC Certificate

Full Quality Assurance System
Directive 98/79/EC on In Vitro Diagnostic Medical Devices,
Annex IV excluding (4, 6)

Registration No.: HL 1804147-1

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

Gamma N-HANCE

Capture-CMV Controls (Kit):

Capture-CMV Positive Control Serum (Weak)
Capture-CMV Negative Control Serum

Capture-R Ready Indicator Red Cells
Capture LISS

Capture-R Controls (Kit):

Capture-R Positive Control Serum (Weak)
Capture-R Negative Control Serum

Bovine Albumin Solution 22%
ImmuAdd
Anti-Fya (Monoclonal) Gamma-clone®
Anti-Fyb (Monoclonal) Gamma-clone®

Report No.: 1111389-10

Effective date: 2020-12-17

Expiry date: 2025-05-26

Issue date: 2022-05-10



Wenxiang Zhang
TUV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 98/79/EC concerning in vitro diagnostic medical devices with the identification number 0197.