EC Design-Examination Certificate

TÜVRheinland®

Directive 98/79/EC on In Vitro Diagnostic Medical Devices,
Annex IV (4)

Registration No.: IL 1804147-1

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

Products: Control reagent for quality control of blood bank reagents by automated methods

- WB corQC

Replaces Certifiicate, Registration No.: IL 60139613 0001

The Notified Body hereby declares that an examination of the design dossier relating to the listed products has been performed according to Annex IV, section 4 of the directive 98/79/EC and that the design of the devices conforms to the requirements of the abovementioned directive.

Report No.: 1104888-10

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TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

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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.

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