

EC Design-Examination Certificate
Directive 98/79/EC Annex IV, Section 4
In Vitro Diagnostic Medical Devices

Registration No.: IL 60139613 0001

Report No.: 60239282 001

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

Product

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

- WB corQC

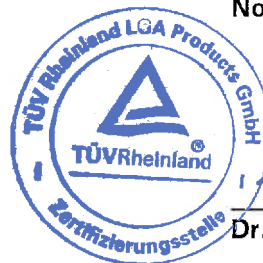
The Notified Body hereby declares that an examination of the design dossier relating to the listed products has been performed according to Annex IV, section 4 of the directive 98/79/EC and that the design of the devices conforms to the requirements of the abovementioned directive.

Expiry Date: 2022-04-10

Effective Date: 2019-05-29

Date: 2019-05-28

Notified Body




Dr. H. Lüdemann

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg

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.