

EC Certificate



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

Registration No.: HL 1381718-1

Manufacturer: Imusyn GmbH & Co.KG
Feodor-Lynen-Str. 5
30625 Hannover
Germany

Products: - HLA SBT Typing Reagents HLA-A, -B, -DR
- recombinant Blood Group Antigen (rBGA) Reagents
- DaraEx plus

Replaces Certificate, Registration No.: HL 60118554 0001

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.

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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.