



Benannt durch/Designated by  
Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln und  
Medizinprodukten  
www.zlg.de  
ZLG-BS-245.10.07



Product Service

# EC Certificate

EC Design-Examination Certificate

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

**No. V7 040330 0183 Rev. 01**

**Manufacturer: DiaMed GmbH**

Pra Rond 23  
1785 Cressier FR  
SWITZERLAND

**Product: Reagents and reagent products for blood typing**

**Model(s): IH-QC Modular System**

Parameters:	Id-n°:	Product Name:	Catalog REF:	Packaging Size:
	08710	IH-QC1	009321	4 x 6 mL
	08720	IH-QC2	009322	4 x 6 mL
	08730	IH-QC3	009323	4 x 6 mL
	08740	IH-QC4	009324	4 x 6 mL
	08750	IH-QC5	009325	4 x 6 mL
	08760	IH-QC6	009326	1 x 6 mL

The Certification Body of TÜV SÜD Product Service GmbH declares that a design examination has been carried out on the respective devices in accordance with IVDD Annex IV (4). The design of the devices conforms to the requirements of this Directive. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:V7\\_040330\\_0183\\_Rev\\_01](http://www.tuvsud.com/ps-cert?q=cert:V7_040330_0183_Rev_01)

**Report No.:** 713249257-02\_SCN

**Valid from:** 2022-04-13

**Valid until:** 2025-05-26

**Date,** 2022-04-13

Christoph Dicks  
Head of Certification/Notified Body