ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 0174 Rev. 01

Manufacturer: **DiaMed GmbH**
Pra Rond 23
1785 Cressier FR
SWITZERLAND

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

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 0174 Rev. 01

Report No.: 713249257-02_SCN

Valid from: 2022-04-13
Valid until: 2024-10-24

Date, 2022-04-13

Christoph Dicks
Head of Certification/Notified Body

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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No. V7 040330 0174 Rev. 01

Model(s): **DiaMed-ID Micro Typing System ID-Cards (monoclonal Reagents)**

Facility(ies): DiaMed GmbH
Pra Rond 23, 1785 Cressier FR, SWITZERLAND

Parameters:

Id-n°:	Product-Name	Catalog REF:	Packaging Size:
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50093	DiaClon ABO/D + Reverse Grouping for Patients	001264	4 x 12 cards
		001267	24 x 12 cards
		001266	60 x 12 cards
		001265	112 x 12 cards
50093	DiaClon ABO/D+ reverse grouping for patients	001265VC	112 x 12 cards
50742	DiaClon ABO/D + Reverse Grouping	001248	60 x 12 cards
		001249	112 x 12 cards

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