Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 037875 0044 Rev. 01

Manufacturer:

KD Medical GmbH
Hospital Products
Charlottenstrasse 65
10117 Berlin
GERMANY

Product Category(ies): **Infusion Sets, Transfusion Sets, Scalp Vein Sets, Syringes, Syringes Insulin, Needles hypodermic, Needles spinal, IV Cannulae with and without safety feature, safety needles (hypodermic needles with needle protection device), Stopcocks, Extension Tubing, Blood lancets, Obturators, Pen needles, Blood Bags, Needles epidural, Syringes with re-use prevention feature with needle (RUP syringes with needle)**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: 713179316

Valid from: 2020-04-20
Valid until: 2024-05-26

Date, 2020-04-20

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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A4 / 07.17