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Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
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Product Service

EC Certificate

Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 037875 0045 Rev. 00

Manufacturer

KD Medical GmbH
Hospital Products
Charlottenstrasse 65
10117 Berlin
GERMANY

Product Category(ies):

**Syringes Catheters, Urine Bags,
Urine Bags Paediatric, Luer Caps,
Injection Stoppers, Spikes,
Single-Use Cannula (blunt),
Syringes with Re-Use Prevention Feature
Without Needle (RUP Syringes Without Needle)**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report No.: 713165011

Valid from: 2020-03-02

Valid until: 2024-05-26

Date, 2020-03-02

Christoph Dicks
Head of Certification/Notified Body

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