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

TÜV SÜD 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 010393 0533 Rev. 02

Manufacturer:

CODAN Medizinische Geräte GmbH & Co KG
Stig Husted-Andersen Strasse 11
23738 Lensahn
GERMANY

Product Category(ies): **Sterile and Non-Sterile Single-Use Medical Devices for Infusion and Transfusion Therapy, Irrigation Sets for Urology and Gynaecology as well as Mixing and Withdraw Products**

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.: 713169069

Valid from: 2020-03-13
Valid until: 2024-05-26

Date, 2020-03-13

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

TÜV®