



Benannt durch/Designated by  
Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln und  
Medizinprodukten  
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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 010393 0533 Rev. 01**

**Manufacturer:**

**CODAN Medizinische Geräte  
GmbH & Co KG**

Stig Husted-Andersen Strasse 11  
23738 Lensahn  
GERMANY

**Facility(ies):**

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

CODAN PORTUGAL Instrumentos Médicos, S.A.  
Rua Stig Husted-Andersen, 4, 2675-492 Odivelas, PORTUGAL

**Product Category(ies): Sterile and Non-Sterile Single-Use Medical  
Devices for Infusion and Transfusion  
Therapy 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.:** 713169066

**Valid from:** 2019-12-09

**Valid until:** 2022-09-02

**Date,** 2019-12-09

Christoph Dicks  
Head of Certification/Notified Body

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ CERTIFICADO ◆ CERTIFICAT