



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

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 074735 0084 Rev. 00

Manufacturer

Covidien llc

15 Hampshire Street
Mansfield MA 02048
USA

Product Category(ies):

**Medical Devices for Wound Care, Wound Closure,
Cardio Thoracic, Respiratory, Blood Collection,
Suction, Aspiration, Gastro-Intestinal Management,
Urology, Enteral Feeding, Operating Room
Support, Needles/Syringes,
and Compression Devices**

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.:

713176602

Valid from:

2020-03-05

Valid until:

2024-05-26

Date,

2020-03-05

Christoph Dicks
Head of Certification/Notified Body