



## **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 074735 0083 Rev. 00

Manufacturer: Covidien IIc

15 Hampshire Street Mansfield MA 02048

**USA** 

Product Category(ies): Medical Devices for Wound Care, Wound Closure.

Cardio-Thoracic, Cardio-Thoracic Drainige.

Respiratory, Suction, Aspiration, Blood Collection.

Gastro-Intestinal Management, Urology,

Vascular Access, Enteral Feeding, Enteral Access, Needles, Syringes, Thermometry, Maternity Care,

Foley Catheters, Foley Catheter Trays,

O.R. Suction, Surgical Suction, Oxygen Delivery, Urine Meter, and Vascular Compression Devices

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.: 713175516

Valid from: 2020-02-28 Valid until: 2024-05-26

Date. 2020-02-28

Christoph Dicks

Head of Certification/Notified Body

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TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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