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Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
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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 077790 0060 Rev. 00

Manufacturer:

Covidien LLC

15 Hampshire Street
Mansfield MA 02048
USA

**Product Category(ies): Oximetry and Capnography Monitor Systems
Temperature Monitor Systems, Patient Warming
Device Systems, Disposable Airway Management
Devices, Tracheal Tubes, Tracheostomy Tubes,
Speaking Valves, and Intubating Stylets, Ventilator
Systems and Patient Interface Circuit Systems,
EEG Monitoring Systems, Breathing Therapy and
Humidification, Heated Inspiratory Line
Humidifiers, Multi-patient Physiologic Monitoring
System and Data Analytics Software,
Gastrointestinal Measurement and Dilation System,
Electrosurgical Diathermy System Electrode.**

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.: 72145607

Valid from: 2020-06-29

Valid until: 2024-05-26

Date, 2020-06-29

Christoph Dicks
Head of Certification/Notified Body