ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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

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 061585 0032 Rev. 00

Manufacturer:

B. Braun Medical AG
Seesatz 17
6204 Sempach
SWITZERLAND

Facility(ies):

B. Braun Medical AG
Route de Sorge 9, 1023 Crissier, SWITZERLAND

B. Braun Medical AG
Seesatz 17, 6204 Sempach, SWITZERLAND

Product Category(ies):

Solutions and powders for disinfection of dental and surgical instruments and devices, endoscopes, anaesthetic equipment, hemodialysis monitors and for use in ultrasonic bathes;
Wipes for cleaning and disinfection of medical devices as ultrasonic probe heads.
Solutions and tissues for surface disinfection of medical equipment, e.g. operating accessories, hospital beds and treatment chairs.
Sterile urinary-catheter-irrigation solutions

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.: 713160957

Valid from: 2019-05-24
Valid until: 2024-05-17

Date, 2019-05-24

Stefan Preiß
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

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