



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 025701 0090 Rev. 01

Manufacturer:

B. Braun Surgical S.A.

Ctra. de Terrassa, 121
08191 Rubi (Barcelona)
SPAIN

Product Category(ies): Sterile absorbable and non absorbable sutures and microsutures with and without needles, combined and non combined with medicinal substances and accessories, Sterile surgical tapes, Sterile surgical meshes combined and non combined with medicinal substances, Surgical stapler, Local haemostatic agents, Absorbable implants for filling of bone defects, Tissue adhesives and accessories, Wound Care Products for Vacuum Therapy, Sterile Suture Procedure packs.

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.: 713165649

Valid from: 2020-03-18

Valid until: 2024-04-26

Date, 2020-03-18

Christoph Dicks
Head of Certification/Notified Body



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