



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 13 11 25701 046

**Manufacturer:** **B. Braun Surgical S.A.**  
Ctra. de Terrassa, 121  
08191 Rubi (Barcelona)  
SPAIN

**Facility(ies):** B. Braun Surgical S.A.  
Ctra. de Terrassa, 121, 08191 Rubi (Barcelona), SPAIN

**Product Category(ies):** **Implants, Suture Material, Tissue Adhesive, local Haemostatics and Sterile Sets/Packs for Sterile Procedure and Surgical Accessories:**

- Microsutures
- Sterile Surgical Sutures
- Sterile Surgical Tapes
- Sterile Surgical Meshes
- Sterile Sets and Packs for Sterile Procedures
- Tissue Adhesives
- Local Hemostatics
- Implants
- Surgical Accessories

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.:** 713033468

**Valid from:** 2014-04-27

**Valid until:** 2019-04-26

**Date,** 2014-04-01

Hans-Heiner Junker



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

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