



Product Service

EC - CERTIFICATE

Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 10 01 49044 009

Manufacturer:	HEBUmedical GmbH Badstraße 8 78532 Tuttlingen GERMANY
Facility(ies):	HEBUmedical GmbH Badstraße 8, 78532 Tuttlingen, GERMANY
Product Category(ies):	Implants for Osteosynthesis, HF-Surgery Devices, HF-Instruments, Single use Skin-Stapler (steril)

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 products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

Report No.: 71358457

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Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

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