



Product Service

EC-CERTIFICATE

Full Quality Assurance System

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

No. G1 09 12 54538 006

Manufacturer: Hermann Medizintechnik GmbHWürttembergische Straße 26
78567 Fridingen/Tuttlingen
GERMANY**Facility(ies):**Hermann Medizintechnik GmbH
Württembergische Straße 26, 78567 Fridingen/Tuttlingen,
GERMANY**Product
Category(ies):****Class IIa:**
Instruments and accessories for invasive and
minimally-invasive surgery for
urology, hysteroscopy, laparoscopy and arthroscopy
Endoscopy units (insufflators, suction-irrigation-units,
arthroscopy pumps, arthroscopy shaver)
Class IIb:
HF-generators and instruments for high frequency
surgery, incl. bipolar instruments
Titanium implant systems: Maxillo-facial, hand and
radius, mandibular, dental, IMF
Steel implants for osteosynthesis

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.

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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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