



Product Service

EC-CERTIFICATE

Full Quality Assurance System

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

No. G1 08 11 19452 013

Manufacturer:	Dufner Instrumente GmbH, Medizintechnik Föhrenstrasse 9 78532 Tuttlingen GERMANY
Facility(ies):	Dufner Instrumente GmbH, Medizintechnik Föhrenstrasse 9, 78532 Tuttlingen, GERMANY
Product Category(ies):	Endoscopes, Endoscope Shafts, High Frequency Instruments, Suction-Irrigation Canullas, -Valves, -Tubes, Insufflation Canullas, Insufflation Devices, Suction- and Irrigation Devices, Sore Clips, Blood Vessels Stripper, Saws, Syringes, Trocar Sleeves

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.: 71345052

Valid until: 2014-01-13

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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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