Product Service

EC-CERTIFICATE

Full Quality Assurance System

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

No. G1 07 11 46083 007

Manufacturer:

MSI MedServ International Deutschland GmbH
Escad Strasse 3
88630 Pfullendorf
GERMANY

Facility(ies):

MSI MedServ International Deutschland GmbH
Escad Strasse 3, 88630 Pfullendorf, GERMANY

Product Category(ies):

Rigid and flexible endoscopes, endoscopes accessories, instruments and lightsources

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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Valid until: 2012-07-03

Date, 2007-12-22

Reiner Krumme

325709

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