



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 48132 005

Manufacturer: LMT
Lammers Medical Technology GmbH
Osterweide 8
23562 Lübeck
GERMANY

Facility(ies): LMT Lammers Medical Technology GmbH
Osterweide 8, 23562 Lübeck, GERMANY

Product Category(ies): MR-compatible incubators,
MR-coils and MR-compatible pressure regulators,
MR-compatible patient monitoring sensors

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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Reiner Krumme



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

Page 1 of 1