



EC Certificate

Full Quality Assurance System Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4) (Devices in Class IIa, IIb or III)

No. G1 046082 0016 Rev. 02

Manufacturer:

Dimeda Instrumente GmbH

Gänsäcker 54 58 78532 Tuttlingen **GERMANY**

Product Category(ies): HF-Instruments and HF-Electrodes,

Wires, Nails, Plates and Screws,

Rib Spreader, Suction / Irrigation Instruments

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 devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.:

713166577

Valid from:

2020-07-02

Valid until:

2024-05-26

Date.

2020-05-04

Christoph Dicks

Head of Certification/Notified Body