



Product Service

# 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 15 03 46082 010

**Manufacturer:** **Dimeda Instrumente GmbH**

Gänsäcker 54+58  
78532 Tuttlingen  
GERMANY

**Facility(ies):**

Dimeda Instrumente GmbH  
Gänsäcker 54+58, 78532 Tuttlingen, GERMANY

**Product  
Category(ies):**

HF-Instruments and HF-Electrodes, HF-Generators, Optics  
(Endoscopes), Rib Spreaders, Orthopedic Implants,  
Wires, Nails, Plates and Screws, Drivers for Surgical  
Instruments, Burrs, Reamers and Drills, Saws and Saw  
Blades, Trocars, Insufflators and Tubing, Suction  
Irrigators, Pumps, Cannulas for Injection and Puncture,  
Veress Needles, Sterile Single use Scalpels and Knives

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.:** 713056182

**Valid from:** 2015-07-02

**Valid until:** 2020-07-01

**Date,** 2015-07-03

Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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