ZERTIFIKAT ◆ CERTIFICATE ◆ 認証証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 14 06 33623 014

Manufacturer:

mahe medical gmbh
Friedrich-Wöhler-Straße 10
78576 Emmingen-Liptingen
GERMANY

Facility(ies):

mahe medical gmbh
Friedrich-Wöhler-Straße 10, 78576 Emmingen-Liptingen,
GERMANY

Product Category(ies):

HF Instruments and –Electrodes, Optics (Endoscopes), Insufflators and tubing, Suction Irrigators, Shaver System, Micromotor Drivers, Kyphoplasty System, Injection-/Puncture needles/Veress Needles, Trocars, Orthopaedic burrs, drills and taps, Orthopaedic reamers and coupling, Rib Spreaders, External Fixation Systems, Cages, Orthopaedic implants, Bone Nails, Bone Plates, Cranioplasty Screws for plates, Bone screws, Bone staples, Bone wires, Blades for bone saws

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.: 713040052

Valid from: 2014-08-01
Valid until: 2019-07-31

Date, 2014-07-15

Hans-Heiner Junker

588415

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