



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 033623 0017 Rev. 01

Manufacturer:

mahe medical gmbh

Friedrich-Wöhler-Straße 10
78576 Emmingen-Liptingen
GERMANY

Product Category(ies):

- HF-Instruments and Electrodes
- Optics (Endoscopes/Laparoscopes/Arthroscopes)
- Suction Irrigators / Surgical Lavage Unit
- Micromotor-Drivers (Surgical Power Tool System)
- Orthopaedic and Laparoscopic Trocars
- Orthopaedic Drills and Taps
- Orthopaedic Reamers
- Saw Blades
- Orthopaedic Internal Fixation Systems, Spinal
(Cervical Plates/Spinal Fusion Cages)
- Orthopaedic Implants
 - Bone Nails
 - Bone Plates
 - Bone Screws
 - Staples
 - Bone Wires
 - Screws for Cranioplasty Plates
 - Craniofacial Fixation Plates

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. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G10336230017Rev.01

Report No.: 713182113

Valid from: 2021-05-20
Valid until: 2024-05-26

Date, 2021-05-20

Christoph Dicks
Head of Certification/Notified Body