



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 16 04 87541 007**

Manufacturer: **MAT GmbH & Co. KG**

Friedrich-Wöhler-Str. 10
78576 Emmingen-Liptingen
GERMANY



Facility(ies):

MAT GmbH & Co. KG
Friedrich-Wöhler-Str. 10, 78576 Emmingen-Liptingen, GERMANY

**Product
Category(ies):**

**Micromotor-Drive, orthopedic burs, taps,
reamers and couplings, saw blades,
external fixation systems,
orthopedic implants: bone nails, bone plates,
bone screws, bone staples and bone wires**

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.

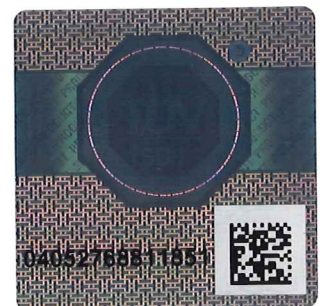
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Date, 2016-05-11

Stefan Preiß



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

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