Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

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 049319 0002 Rev. 00

Manufacturer:

joimax® GmbH
Amalienbadstraße 41
RaumFabrik 61
76227 Karlsruhe
GERMANY

Facility(ies):

joimax® GmbH
Amalienbadstraße 41, RaumFabrik 61, 76227 Karlsruhe,
GERMANY

Product Category(ies):

Instruments, devices, accessories and corresponding disposables for endoscopic spine surgery: endoscopes, irrigation pumps with corresponding tubing set, shrill systems and shrill blades, sets for spine surgery, reamers, drills, needles, HF/RF generators and probes, cages, pedicle screws and rods.

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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Valid from: 2018-12-24
Valid until: 2023-12-23

Date, 2018-12-21

Stefan Preiß

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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany