



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 03 44166 023

Manufacturer: **GIMMI GmbH**
Carl-Zeiss-Strasse 6
78532 Tuttlingen
GERMANY

Facility(ies): GIMMI GmbH
Carl-Zeiss-Strasse 6, 78532 Tuttlingen, GERMANY

Gimmi GmbH
Lohmehlenring 90, 78532 Tuttlingen, GERMANY

Product Category(ies): **HF Instruments and accessories,
HF units, endoscopic systems and accessories,
insufflators, surgical instruments (class IIa),
orthopedic implants,
aneurysm clips (class III),
arthro-perfusion pump,
shaver system, shaver blades**

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.

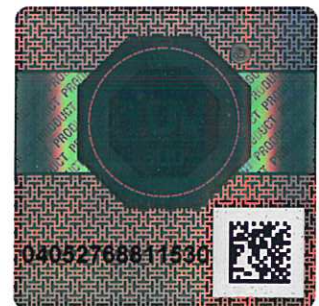
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Date, 2016-04-08

Stefan Preiß



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

Page 1 of 1