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Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
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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 020326 0060 Rev. 02

Manufacturer: **Institut Straumann AG**
Peter Merian-Weg 12
4002 Basel
SWITZERLAND

Product Category(ies): **Implants, dental materials, prosthetic components and their systems/instruments for use in oral and extraoral implantology (maxillofacial surgery) and dental prosthetic restoration.**

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.: 713181490

Valid from: 2020-03-24

Valid until: 2024-05-26

Date, 2020-03-24

Christoph Dicks
Head of Certification/Notified Body

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ CERTIFICADO ◆ CERTIFICAT