



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 11 02 20326 035

**Manufacturer:****Institut Straumann AG**

Peter Merian-Weg 12  
4002 Basel  
SWITZERLAND

**Facility(ies):**

Institut Straumann AG  
Peter Merian-Weg 12, 4002 Basel, SWITZERLAND

Straumann CAD/CAM GmbH  
Lochamer Schlag 6, 82166 Gräfelfing, GERMANY

Straumann CAD/CAM GmbH  
Annaberger Straße 240, 09125 Chemnitz, GERMANY

Straumann CAD/CAM GmbH  
Rudower Chaussee 29, 12489 Berlin, GERMANY

**Product  
Category(ies):**

**Implants, regenerative materials, prosthetic components,  
medical software and their systems/instruments for use in  
oral and extraoral implantology (maxillo-facial 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.:**

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**Valid from:**

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**Valid until:**

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**Date,** 2011-03-11

Hans-Heiner Junker

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