



Product Service

EC Design Examination Certificate

(Annex II, section 4 of the Directive 93/42/EEC on Medical Devices)

No. G7 10 09 20326 029

Manufacturer: Institut Straumann AG

Peter-Merian-Weg 12
4002 Basel
SWITZERLAND

Product: Non-Active Implants

Biodegradable, synthetic hydrogel membrane
for aid in oral guided bone regeneration

Model(s): Straumann® MembraGel

The Certification Body of TÜV SÜD Product Service GmbH declares that a design examination has been carried out on the aforementioned devices according to Annex II, section 4 of the Directive 93/42/EEC on Medical Devices. The design of the devices conforms to the provisions of this Directive. For marketing of these products an additional Annex II.3 certificate is mandatory. See also notes overleaf.

Report No.: 71376278

Valid until: 2015-09-22

Date, 2010-10-01

H.-H.
Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

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