



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 12 07 43306 095

Manufacturer: Ivoclar Vivadent AG

Bendererstraße 2
9494 Schaan
LIECHTENSTEIN

**Product
Category(ies):**

**Dental materials: Filling materials
(including sealants, liners, base materials,
temporary materials), cements, adhesives, desensitizers
etching materials, bleaching materials,
root canal posts, polishing instruments,
resin and porcelain teeth, denture base materials,
materials for inlays, onlays, crowns and bridges,
non-precious dental alloys,
antibacterial protective varnish for dental use,
dental abutments, dental curing lights,
dental instruments**

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.: 713007129

Valid from: 2012-10-01

Valid until: 2014-11-30

Date, 2012-10-04

Hans-Heiner Junker



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

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Facility(ies):

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