



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 14 11 78535 025

Manufacturer:**3M Deutschland GmbH**

Carl-Schurz-Straße 1
41453 Neuss
GERMANY

**Product
Category(ies):**

Implant materials for dentistry as filling materials as well as crown and bridge materials, abutments, luting cements, adhesives, etching agents, disinfectants and micro-blasters, dental materials for surface preparation and endodontic points, rotary instruments, caries indicators

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.:

713051701

Valid from:

2015-01-21

Valid until:

2020-01-19

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Hans-Heiner Junker



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

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Facility(ies):

3M Deutschland GmbH
ESPE Platz, 82229 Seefeld, GERMANY

3M Deutschland GmbH
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