Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

TÜV SÜD 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 078535 0039 Rev. 00

Manufacturer:

3M Deutschland GmbH
Carl-Schurz-Straße 1
41453 Neuss
GERMANY

Facility(ies):

3M Deutschland GmbH
ESPE Platz, 82229 Seefeld, GERMANY

3M Deutschland GmbH
Ohmstraße 3, 86899 Landsberg, GERMANY

Product Category(ies):

Implant materials for dentistry as filling materials as well as crown and bridge materials, luting cements, adhesives, etching agents and micro-blasters, dental materials for surface preparation and endodontic posts with corresponding drills

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.: 713158370

Valid from: 2019-09-25
Valid until: 2024-05-26

Date, 2019-09-25

Stefan Preiß
Head of Certification/Notified Body

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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany