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 015409 0035 Rev. 00

Manufacturer:

VDW GmbH
Bayerwaldstraße 15
81737 München
GERMANY

Facility(ies):

VDW GmbH
Bayerwaldstraße 15, 81737 München, GERMANY

Product Category(ies):

Dental instruments, Devices for root canal treatment and accessories for the product(s)/product category(ies):
- Lubricant for endodontic instruments in the root canal preparation
- Rotating root canal instruments
- Root canal filling points
- Paper points to dry the root canal
- Sterile root canal instruments for single use
- Device for root canal length determination
- Root canal posts
- Electronic powered motor for root canal root instruments
- Endo contra angle
- Software as accessory for endodontic equipment

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.: 713145437

Valid from: 2019-06-07
Valid until: 2024-05-26

Date, 2019-06-07

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

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