

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 15 08 15409 023

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

(see attachment)

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.:

713067667

Valid from: Valid until:

2016-01-16 2021-01-15

Date, 2015-12-22

Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123 Page 1 of 2