



Product Service

# EC Certificate

## Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V  
(Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 15 08 15409 024

**Manufacturer:** VDW GmbH  
Bayerwaldstraße 15  
81737 München  
GERMANY



**Facility(ies):** VDW GmbH  
Bayerwaldstraße 15, 81737 München, GERMANY

**Product Category(ies):** Sterile root canal instruments for repeated use

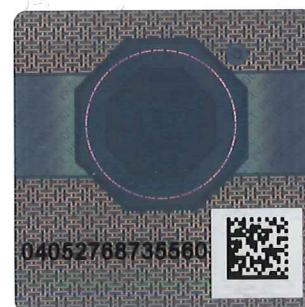
The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

**Report No.:** 713067667

**Valid from:** 2016-01-16  
**Valid until:** 2021-01-15

**Date,** 2015-12-21

Hans-Heiner Junker



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

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