



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 015409 0027 Rev. 00

Manufacturer

VDW GmbH

Bayerwaldstraße 15 81737 München GERMANY

Facility(ies):

Category(ies):

VDW GmbH

Bayerwaldstraße 15, 81737 München, GERMANY

Product

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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.:

713145440

Valid from: Valid until:

2019-06-07 2024-05-26

Date,

2019-06-07

Stefan Preiß

1. Punil

Head of Certification/Notified Body

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