



## **EC** Certificate

Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)

No. G2 051317 0005 Rev. 01

Manufacturer: MK-dent GmbH

Marie-Curie-Straße 2 22941 Bargteheide GERMANY

Facility(ies): MK-dent GmbH

Marie-Curie-Straße 2, 22941 Bargteheide, GERMANY

Product Turbines, high and low speed handpieces, prophylaxis handpieces, motors and couplings

including spare parts and accessories for the

dental area

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.: 713162758

 Valid from:
 2020-01-24

 Valid until:
 2024-05-26

Date, 2020-01-24

Christoph Dicks

Head of Certification/Notified Body

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

