



Benannt durch/Designated by
 Zentralstelle der Länder
 für Gesundheitsschutz
 bei Arzneimitteln und
 Medizinprodukten
 www.zlg.de
 ZLG-BS-244.10.08



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 045041 0025 Rev. 00

Manufacturer: **Maxtec**
 2305 South 1070 West
 Salt Lake City UT 84119
 USA

**Product Category(ies): Electrochemical Oxygen Sensors,
 Analyzers, Monitors, Air/Oxygen
 Blenders, Flowmeters and
 CO2 Indicators**

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. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:G10450410025 Rev. 00](http://www.tuvsud.com/ps-cert?q=cert:G10450410025Rev.00)

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Valid from: 2021-03-12

Valid until: 2024-05-26

Date, 2021-03-12

Christoph Dicks
 Head of Certification/Notified Body