



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 11 09 45041 012

Manufacturer: **Maxtec**
6526 So. Cottonwood St. (300W)
Salt Lake City UT 84107
USA

EC-Representative: **QNET BV**
Hommerterweg 286
6436 AM Amstenrade
THE NETHERLANDS

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

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.: DM1108916

Valid from: 2011-12-19
Valid until: 2016-12-18



Date, 2011-11-21

Hans-Heiner Junker

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

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Facility(ies):

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