



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 13 03 66456 009

Manufacturer: **Magnifico Healthcare, Inc**
1351 S. Leavitt Av # 103B
Orange City, FL 32763
USA

EC-Representative: **Geranswers**
Rua Jos Rodrigues Miguis n141
2870-466 Montijo
PORTUGAL

Product Category(ies): **Temperature Probe for Medical Use, SpO2 Sensor, Ultrasonic Fetal Heart Detector, Oximeter, Electrocardiograph, Disposable Blood Pressure Transducer, Oxygen Sensor for Medical Use**

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.: 748467106

Valid from: 2013-06-09

Valid until: 2018-06-08

Date, 2013-05-06

Hans-Heiner Junker



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

Page 1 of 2



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