



Product Service

EC - CERTIFICATE

Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 08 07 24497 019

Manufacturer: **NONIN MEDICAL, INC.**
 13700 1st Avenue North
 Plymouth MN 55441-5443
 USA

EC-Representative: **MPS Medical Product Service GmbH**
 Borngasse 20
 35619 Braunfels
 GERMANY

Product Category(ies): **Pulse Oximeters, Breathing Monitors, Non-Invasive Blood Pressure Monitors and Sterile Sensors**

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 products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

Report No.: NM805556

Valid until: 2013-12-01

Date, 2008-12-02

Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.



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Facility(ies):

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