



Product Service

EC - CERTIFICATE

Full Quality Assurance System

(Annex 2, section 3 of the Directive 90/385/EEC on Active Implantable Medical Devices)

No. I1 09 09 19742 039

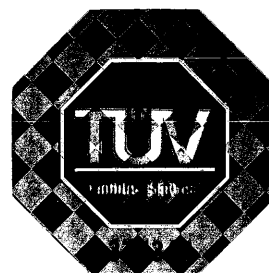
Manufacturer: **VascoMed GmbH**
Hertzallee 1
79589 Binzen
GERMANY

Product: **Accessories for Implantable Leads for AIMDs**

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 families according to Annex 2, section 3 of the Directive 90/385/EEC on Active Implantable Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of the product an additional Annex 2.4 certificate is mandatory. See also notes overleaf.

Report No.: 71352991

Valid until: 2014-09-01



Date, 2009-09-02

Hans-Heiner Junker

TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 90/385/EEC concerning Active Implantable Medical Devices with identification no. 0123.

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Facility(ies):

VascoMed GmbH
Hertzallee 1, 79589 Binzen, GERMANY

Design

Facility(ies):

VascoMed GmbH
Hertzallee 1, 79589 Binzen, GERMANY