



Product Service

EC - CERTIFICATE

Full Quality Assurance System

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

No. G1 09 09 19742 041

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

Product Category(ies): **Ablation Catheters,
Temporary Catheters and Electrodes for
Stimulation and Electrophysiology,
Vascular Catheters, Heartwires,
Accessories for Catheters and Electrodes,
Introducer Sheaths and Sets**

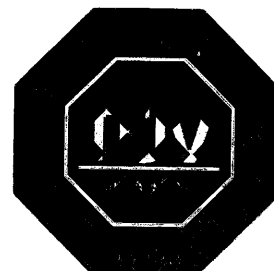
The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical 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.: 71352991

Valid until: 2014-09-01

Date, 2009-10-09

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):

VascoMed GmbH
Hertzallee 1, 79589 Binzen, GERMANY

VascoMed GmbH
Marie-Curie-Strasse 8, 79539 Lörrach, GERMANY

Design

Facility(ies):

VascoMed GmbH
Hertzallee 1, 79589 Binzen, GERMANY