ZERTIFIKAT ◆ CERTIFICATE ◆ 認証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 14 08 19742 059

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 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.:	713047033
Valid from:	2014-09-02
Valid until:	2019-09-01

Date, 2014-09-01

Hans-Heiner Junker

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TÜV SÜD Product Service GmbH is Notified Body 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

VascoMed GmbH
Pasteurallee 1, 79589 Binzen, GERMANY

Design Facility(ies):

VascoMed GmbH
Hertzallee 1, 79589 Binzen, GERMANY

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