Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

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 019742 0092 Rev. 00

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

Product Category(ies): **Ablation Catheters, Temporary Catheters and Electrodes for Stimulation and Electrophysiology Diagnostics, Accessories for Catheters and Electrodes**

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.: 713161167

Valid from: 2019-08-27
Valid until: 2024-05-26

Date, 2019-08-27

Stefan Preiß
Head of Certification/Notified Body

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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT
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No. G1 019742 0092 Rev. 00

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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