

EU QUALITY MANAGEMENT SYSTEM CERTIFICATE

Certificate no.
20373GB448260602

Final Assessment Report no.
20373AU04F

Effective date
2026-06-02

Expiry date
2028-03-14

This is to certify that the quality system of

VascoMed GmbH

Hertzallee 1, 79589 Binzen, Germany

SRN: DE-MF-000005571

For design & development, manufacturing and final product inspection/testing of
Medical devices/groups of medical devices at locations as listed on the following pages

has been assessed and found to comply with respect to

**The conformity assessment procedure described in Annex IX,
Chapters I and III of Regulation (EU) 2017/745 on Medical Devices**

Any applicable limitations for certain medical devices are included in the following list or recorded in the final assessment report. This certification is subject to surveillance by DNV MEDCERT.

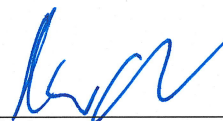
Place and date
Hamburg, 2026-06-02

For the issuing office
DNV MEDCERT GmbH – Notified Body 0482
Brooktorkai 18, 20457 Hamburg, Germany



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

BS-MDR-096



Marcus Harder
Certification Body Operations

The certificate is only valid when provided entirely with all of its pages. To verify the validity of this certificate, contact medcert-info@dnv.com



Certificate no.: [20373GB448260602](#)
Place and date: [Hamburg, 2026-06-02](#)

Sites covered by this certificate

VascoMed GmbH, Hertzallee 1, 79589 Binzen, Germany

VascoMed GmbH, Eulerstrasse 1, 79589 Binzen, Germany

Preceding certificate

Certificate no.	Issue date	Identification of changes
20373GB448250402	2025-04-02	Change of warehouse address according to WO-015099





DNV

Certificate no.: [20373GB448260602](#)
Place and date: [Hamburg, 2026-06-02](#)

Products covered by this certificate

Class III medical devices

For placing on the market of class III medical devices covered by this certificate, an additional EU Technical Documentation Assessment Certificate according to Annex IX Chapter II of Regulation (EU) 2017/745 is required, which also contains the exact determination of medical devices covered by certification.

Category	Medical devices/groups of medical devices
MDN 1203	Non-active non-implantable guide catheters, balloon catheters, guidewires, introducers, filters, and related tools

