



EU Quality Management System Certificate

Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapter I

Certificate No. G15 011634 0256 Rev. 00

Manufacturer:
medela 

Medela AG
Lättichstrasse 4b
6340 Baar
SWITZERLAND

SRN Manufacturer - CH-MF-000018913

Authorized Representative:

Medela Medizintechnik GmbH & Co. Handels KG
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The quality management system has been evaluated in accordance with Regulation (EU) 2017/745, Annex IX Chapter I with a positive result.

Details on devices covered by the quality management system are described on the following page(s). The report referenced below summarises the results of the assessment and includes reference to relevant CS, harmonised standards and test reports.

The certified quality management system is subject to periodical surveillance.

If class I devices in sterile conditions, with measuring function, or reusable surgical instruments are covered by this certificate, the audit was limited to the respective aspects relating to

- establishing, securing, and maintaining sterile conditions,
- conformity of the devices with the metrological requirements,
- reuse of the device, in particular cleaning, disinfection, sterilization, maintenance and functional testing and the related instructions for use.

If class IIa or class IIb devices are covered by this certificate, the quality management system assessment was accompanied by the assessment of technical documentation for devices selected on a representative basis. The periodical surveillance includes further assessment of the technical documentation on the basis of representative samples.

If class III (excluding custom-made implantable devices) or class IIb implantable devices are covered by this certificate, an EU Technical Documentation Assessment Certificate in accordance with Annex IX Chapter II is required before placing them on the market.

All applicable requirements of the Testing, Certification, Validation and Verification Regulations TÜV SÜD Group have to be complied with.

For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G15 011634 0256 Rev. 00

Report No.: 713265779
Valid from: 2026-04-28
Valid until: 2031-04-27

Christoph Dicks
Head of Certification/Notified Body

Issue date: 2026-04-28



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Classification: Class I
Device Group: A06 - DRAINAGE AND FLUIDS COLLECTION DEVICES
Device Properties: MDS 1005 - Devices in sterile condition

Classification: Class I
Device Group: Z12 - INSTRUMENTS FOR FUNCTIONAL EXPLORATIONS AND THERAPEUTIC INTERVENTIONS
Device Properties: MDS 1005 - Devices in sterile condition

The validity of this certificate depends on conditions and/or is limited to the following: - none -

Revision History:

Rev.	Dated	Report	Description
00	2026-04-28	713265779	Initial issuance