



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zfg.de
BS-MDR-099



EU Quality Management System Certificate

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

Certificate No. G15 129096 0004 Rev. 00

Manufacturer:

mylife Diabetes Care AG

Lyssachstrasse 40
3400 Burgdorf
SWITZERLAND

SRN Manufacturer - CH-MF-000044659

Authorized Representative:

mylife Distribution GmbH
Höchster Strasse 70, 65835 Liederbach, GERMANY

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 129096 0004 Rev. 00

Report No.: 713408611

Preceding Certificate No.: G10 129096 0002 Rev. 03

Valid from: 2026-05-18

Valid until: 2030-03-17

Christoph Dicks
Head of Certification/Notified Body

Issue date: 2026-05-18



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Classification: Class IIa
Device Group: MDN 1202 - Non-active non-implantable devices for administration, channelling and removal of substances, including devices for dialysis

Classification: Class IIb
Device Group: A03040101 - KITS FOR CONTINUOUS SUBCUTANEOUS INSULIN INFUSION

Intended Purpose: Infusion Sets are used for the continuous subcutaneous delivery of insulin from the external insulin infusion pump YpsoPump

Classification: Class IIb
Device Group: Z1204021601 - PORTABLE INSULIN INFUSION INSTRUMENTS
Intended Purpose: Insulin Infusion Pumps are intended for continuous subcutaneous insulin delivery

The validity of this certificate depends on conditions and/or is limited to the following: ./.

Revision History:

Rev.	Dated	Report	Description
00	2026-05-18	713408611	Amended: Change of authorized representative Administrative merge / transfer to new Certificate Type