



EU Quality Assurance Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex XI Part A
(Class I Devices in sterile condition, with measuring function or reusable surgical instruments)

No. G21 044934 0010 Rev. 00

Manufacturer:

Intensiv Medical Products IMPROMEDIFORM GmbH

Gielster Stück 11
58513 Lüdenscheid
GERMANY

SRN Manufacturer - DE-MF-000014610

The Certification Body of TÜV SÜD Product Service GmbH certifies that the manufacturer has established, documented and implemented a quality management system as described in Article 10 (9) of the Regulation (EU) 2017/745 on medical devices. Details on device categories covered by the quality management system are described on the following page(s).

The Report referenced below summarises the result of the assessment and includes reference to relevant CS, harmonized standards and test reports. The conformity assessment has been carried out according to Annex XI Part A of this regulation with a positive result.

As applicable the involvement of the notified body is limited to the 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.

The certified quality assurance system is subject to periodical surveillance by TÜV SÜD Product Service GmbH. 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:G21_044934_0010_Rev.00

Report No.: 713333504

Valid from: 2024-08-16

Valid until: 2029-08-15

Issue date: 2024-08-16

Christoph Dicks
Head of Certification/Notified
Body



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Classification: Class I
Device Group: A99 - DEVICES FOR ADMINISTRATION, WITHDRAWAL AND COLLECTION - OTHER
Device Properties: MDS 1005.1 - Ethylene Oxide sterilization

Classification: Class I
Device Group: A0799 - ADAPTERS, CONNECTORS, RAMPS, STOPCOCKS, CAPS - OTHER
Device Properties: MDS 1005.1 - Ethylene Oxide sterilization

Classification: Class I
Device Group: A030499 - ADMINISTRATION KITS - OTHER
Device Properties: MDS 1005.1 - Ethylene Oxide sterilization

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

Revision History:

Rev.	Dated	Report	Description
00	2024-08-16	713333504	Initial issuance