



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



Product Service

## EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III  
(Class IIa and Class IIb Devices)

**No. G10 012974 0611 Rev. 03**

**Manufacturer:**

**B. Braun Melsungen AG**

Carl-Braun-Str. 1  
34212 Melsungen  
GERMANY

**SRN Manufacturer:**

DE-MF-000000201

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 IX Chapter I and III of this regulation with a positive result.

The quality management system assessment was accompanied by the assessment of technical documentation for devices selected on a representative basis.

The certified quality management system is subject to periodical surveillance by TÜV SÜD Product Service GmbH. The surveillance assessment shall also include an assessment of the technical documentation for the device or devices concerned on the basis of further representative samples. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with.

For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:G10 012974 0611 Rev. 03](http://www.tuvsud.com/ps-cert?q=cert:G10 012974 0611 Rev. 03)

**Report No.:**

713225005

**Preceding Certificate No.:**

G10 012974 0611 Rev. 02

**Valid from:**

2022-11-10

**Valid until:**

2025-03-12

**Date of Initial Issuance:**

2020-03-13

Christoph Dicks  
Head of Certification/Notified Body

**Issue date:** 2022-11-10





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**Classification:** IIa  
**Device Group:** A030101 - INFUSION CONTROLLERS  
**Intended Purpose:** -

**Classification:** IIb  
**Device Group:** Z120303 - INFUSION INSTRUMENTS  
**Intended Purpose:** Transportable infusion pump that is used in combination with authorized disposables and accessories.  
The pump is intended for use on adults, pediatrics, and neonates for the intermittent or continuous delivery of parenteral and enteral fluids through clinically accepted routes of administration. These routes include, but are not limited to intravenous, intra-arterial, subcutaneous, epidural, irrigation and enteral. The system is used for the delivery of fluids indicated for infusion therapy.

**Classification:** IIa  
**Device Group:** A020102 - INFUSION AND IRRIGATION SYRINGES, SINGLE-USE  
**Intended Purpose:** -

**Classification:** IIb  
**Device Group:** Z12030382 - INFUSION INSTRUMENTS - SOFTWARE ACCESSORIES  
**Intended Purpose:** Software application platform that is intended to provide bidirectional data communication with authorized medical devices and their accessories. The software application platform is intended to provide gateway functions, visualization of data and configuration of data sets for authorized medical devices and accessories. These data sets include, but are not limited to drug data sets (Drug Library Data) and pump modification data sets (Pump Configuration Data).

**Classification:** IIa  
**Device Group:** A010101 - HYPODERMIC NEEDLES  
**Intended Purpose:** -

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





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### Revision History:

| Rev. | Dated      | Report                      |
|------|------------|-----------------------------|
| 00   | 2020-03-13 | 713169695                   |
| 01   | 2020-11-19 | 713169695                   |
| 02   | 2021-12-28 | 713188740_CN / 713188421_CN |

