



## 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. 13**

**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, 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:G10 012974 0611 Rev. 13](http://www.tuvsud.com/ps-cert?q=cert:G10_012974_0611_Rev.13)

<b>Report No.:</b>	713375453
<b>Preceding Certificate No.:</b>	G10 012974 0611 Rev. 12
<b>Valid from:</b>	2025-06-03
<b>Valid until:</b>	2030-03-12
<b>Date of Initial Issuance:</b>	2020-03-13

**Issue date:** 2025-06-03

Head of Certification/Notified Body



## 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. 13**

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

**Classification:** Class 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.

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

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

**Classification:** Class IIa  
**Device Group:** C010101 - PERIPHERAL I.V. CATHETERS  
**Intended Purpose:** -



## 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. 13**

<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A070199 - ADAPTERS AND CONNECTORS - OTHER
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A040101 - ADMINISTRATION AND ASPIRATION FILTERS
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A070501 - CAPS OR OBTURATORS, NON-PERFORABLE
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A070502 - CAPS OR OBTURATORS, PERFORABLE
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A060101 - VACUUM AND GRAVITY DRAINAGE SYSTEMS
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A018003 - NEEDLE INTRODUCERS
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A010302 - PLEXUS BLOCK NEEDLES AND KITS
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A0703 - STOPCOCKS
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A030103 - ENTERAL FEEDING CONTROLLERS
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A030201 - EXTENSIONS
<b>Intended Purpose:</b>	-



## 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. 13**

<b>Classification:</b>	Class IIa
<b>Device Group:</b>	G020201 - NASOGASTRIC INTESTINAL TUBES
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A070103 - INFUSION LINES ADAPTERS AND CONNECTORS
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A020106 - INSULIN SYRINGES, SINGLE-USE
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIb
<b>Device Group:</b>	A050101 - ELASTOMERIC SYSTEMS - FIXED FLOW
<b>Intended Purpose:</b>	Disposable elastomeric infusion pump system is a non-electrically driven portable infusion device, enabling patients to be treated in an ambulatory manner. The device is indicated for delivering a pre-determined amount of medication to the patient via intravenous, subcutaneous or epidural routes (according to pump model and SPCs of drugs) in a continuous and accurate manner.
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A060201 - EXTERNAL DRAINAGE CATHETERS AND KITS (ABSCESSSES, GALLSTONES, CYSTS)
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A0799 - ADAPTERS, CONNECTORS, RAMPS, STOPCOCKS, CAPS - OTHER
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A060203 - PLEURAL DRAINAGES WITH VALVE AND KITS
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	C010280 - CENTRAL VENOUS CATHETERS - ACCESSORIES
<b>Intended Purpose:</b>	-



## 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. 13**

**Classification:** Class IIa  
**Device Group:** Z120303 - INFUSION INSTRUMENTS  
**Intended Purpose:** -

**Classification:** Class IIb  
**Device Group:** V0999 - FLUIDS/GASES FOR CLINICAL/THERAPEUTICAL USE - OTHER  
**Intended Purpose:** Ready for use, sterile, single use medical device, intended to be used for irrigation applications (irrigation solution)

**Classification:** Class IIb  
**Device Group:** U040102 - KITS WITH DRAINAGE CATHETERS AND INTRODUCERS  
**Intended Purpose:** Suprapubic catheterization of the bladder after surgery, in case of bladder dysfunction, urinary retention, for diagnostic purposes and for urine assessment

**Classification:** Class IIa  
**Device Group:** U040102 - KITS WITH DRAINAGE CATHETERS AND INTRODUCERS  
**Intended Purpose:** -

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

### Revision History:

Rev.	Dated	Report	Description
00	2020-03-13	713169695	-
01	2020-11-19	713169695	-
02	2021-12-28	713188740_CN / 713188421_CN	-
03	2022-11-10	713225005	-
04	2023-03-31	713270133	Supplemented: Device(s)/group of device(s) added
05	2023-05-22	713282403	- Supplemented: Device(s)/group of device(s) added
06	2023-11-10	713309567 / 713309565	Supplemented: Device(s)/group of device(s) added
07	2024-02-15	713279371 / 713313043 / 713316921 / 713316928 / 713316930 / 713316916 / 713316919 / 713316912	Supplemented: Device(s)/group of device(s) added
08	2024-04-23	713332639	Supplemented: Device(s)/group of



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. 13

09	2024-05-28	713308882	device(s) added Supplemented: Device(s)/group of device(s) added
10	2024-09-16	713339665, 713339669, 713339656, 713282405	Supplemented: Device(s)/group of device(s) added
11	2024-11-13	713281980	Supplemented: Device(s)/group of device(s) added
12	2025-03-13	713340251 / 713350098	Renewal of certificate Supplemented: Device(s)/group of device(s) added
13	2025-06-03	713375453	Supplemented: Device(s)/group of device(s) added