

Benannt durch Designated by Zentralstelle der Länder für Gresundheinsschurz bei Arzneimitteln und Medizinprodukten BS-MDR-099





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 010578 0039 Rev. 09

Manufacturer:

Drägerwerk AG & Co. KGaA

Moislinger Allee 53-55 23542 Lübeck GERMANY

SRN Manufacturer:

DE-MF-000005329

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 010578 0039 Rev. 09

Report No.:

713253108_CN

Preceding Certificate No.:

G10 010578 0039 Rev. 08

 Valid from:
 2023-03-14

 Valid until:
 2025-03-17

Date of Initial Issuance: 2020-03-18

Issue date: 2023-03-14

Christoph Dicks Head of Certification/Notified Body



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Classification:	Class Ila	
Device Group:	R02 - BREATHING CIRCUITS AND CATHETER MOUNTS R0301 - RESPIRATORY MASKS R030201 - VENTILATION BALLOONS R0401 - VENTILATION FILTERS R0402 - NATURAL BREATHING FILTERS Z120301 - ANAESTHESIA AND PULMONARY VENTILATION SUPPORT INSTRUMENTS Z120309 - MEDICAL/MEDICINAL GAS PIPELINE SYSTEMS AND RELATED ACCESSORIES	
Intended Purpose:	.	
Classification:	Class IIa	
Device Group:	Z12040192 - GENERAL MEDICINE DIAGNOSIS AND MONITORING INSTRUMENTS - MEDICAL DEVICE SOFTWARE	
Intended Purpose:	-	
Classification:	Class IIa	
Device Group:	Z120302 - VITAL SIGNS MONITORING INSTRUMENTS	
Intended Purpose:	-	
Classification:	Class IIa	
Device Group:	A060304 - INTRA-OPERATION FLUID COLLECTION DEVICES	
Intended Purpose:	-	
Classification:	Class IIb	
Device Group:	Z12040192 - GENERAL MEDICINE DIAGNOSIS AND MONITORING INSTRUMENTS - MEDICAL DEVICE SOFTWARE Software intended to provide clinical information for the purpose of supporting patient management and the decision making process	
Intended Purpose:		
Classification:	Class IIb	
Device Group:	Z120804 - NEONATOLOGY INSTRUMENTS	

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Intended Purpose:	Warming therapy devices intended to provide controlled ambient conditions for premature babies and neonates in closed and open care therapy	
Classification:	Class IIb	
Device Group:	Z120301 - ANAESTHESIA AND PULMONARY VENTILATION SUPPORT INSTRUMENTS Devices for the purpose of ventilation and/or anesthesia	
Intended Purpose:		
Classification:	Class IIb	
Device Group:	Z120309 - MEDICAL/MEDICINAL GAS PIPELINE SYSTEMS AND RELATED ACCESSORIES Devices intended to distribute or supply gases, vacuum, electricity or data to equipment in diagnostic, therapy or surgery	
Intended Purpose:		
Classification:	Class IIb	
Device Group:	R020107 - THERMOREGULATED BREATHING CIRCUITS	
Intended Purpose:	Inspiratory (and expiratory) heated disposable breathing circuit for conducting humidified breathing gas from humidifier to patient	
Classification:	Class IIb	
Device Group:	R020101 - STANDARD BREATHING CIRCUITS	
Intended Purpose:	Devices intended to administer gases for the purpose of ventilation	

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

Revision History:

Rev.	Dated	Report
00	2020-03-18	713169482
01	2021-07-02	713184148
02	2021-09-30	713215188
03	2021-10-01	713215832
04	2021-10-04	713215842
05	2021-10-04	713219421
06	2021-11-22	713229134

Description

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No. G10 010578 0039 Rev. 09

07	2022-02-21	713213004
08	2022-10-06	713225304_CN
09	2023-03-14	713253108_CN

Supplemented: Device(s)/group of device(s) added

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CATHETER MOUNTS AND AIRWAY CONNECTORS

