ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 049861 0162 Rev. 02

Manufacturer:

ResMed Pty Ltd
1 Elizabeth Macarthur Drive
Bella Vista NSW 2153
AUSTRALIA

SRN Manufacturer: **AU-MF-000011753**

Authorized Representative:

ResMed SAS
Parc Technologique de Lyon,
292 Allée Jacques Monod,
69791 Saint Priest Cedex, FRANCE

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 049861 0162 Rev. 02

Report No.: **JA36117609**

Preceding Certificate No.: **G10 049861 0162 Rev. 01**

Valid from: **2022-08-04**
Valid until: **2025-10-06**

Date of Initial Issuance: **2020-10-07**

Christoph Dicks
Head of Certification/Notified Body

Issue date: 2022-08-04

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123
TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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Classification:	IIa
Device Group:	R020107 - THERMOREGULATED BREATHING CIRCUITS
Intended Purpose:	-/-

Classification:	IIa
Device Group:	R020104 - CPAP AND NIV BREATHING CIRCUITS
Intended Purpose:	-/-

Classification:	IIa
Device Group:	Z120301 - ANAESTHESIA AND PULMONARY VENTILATION SUPPORT INSTRUMENTS
Intended Purpose:	-/-

Classification:	IIa
Device Group:	R030101 - VENTILATION MASKS
Intended Purpose:	-/-

Classification:	IIa
Device Group:	R0203 - ANAESTHESIA AND RESUSCITATION CONNECTORS
Intended Purpose:	-/-

Classification:	IIa
Device Group:	R0280 - BREATHING CIRCUITS AND CATHETER MOUNTS - ACCESSORIES
Intended Purpose:	-/-

Classification:	IIa
Device Group:	R030102 - AIR/OXYGEN MASKS AND NASAL CANNULAS

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No. G10 049861 0162 Rev. 02

Intended Purpose:	-/-
Classification:	IIa
Device Group:	R040199 - VENTILATION FILTERS - OTHER
Intended Purpose:	-/-
The validity of this certificate depends on conditions and/or is limited to the following:	-/-

Revision History:

Rev.	Dated	Report
00	2020-10-07	JA1437662
01	2022-02-10	JA1634396

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