Resmed

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Declaration of Conformity

Manufacturer:	Authorized Representative:	Notified Body:
ResMed Corp. 9001 Spectrum Center Blvd. San Diego CA 92123 USA	ResMed SAS Parc Technologique de Lyon 292 Allée Jacques Monod 69791 Saint Priest Cedex France	TÜV SÜD Product Service GmbH Ridlerstraße 65 80339 München Germany

Product: ResScan Essentials

Intended Use:
ResScan Essentials is intended to augment the standard follow-up care of patients by transferring and analyzing machine and therapeutic information from ResMed compatible therapy devices to the ResScan Essentials application. ResScan Essentials is intended to be used by clinicians.

Classification: IIa according to Rule 11

EMDN: Z1203019092 Various Instruments for Anaesthesia and Pulmonary Ventilation Support - Medical Device Software

Conformity Assessment Route: Annex IX (excluding Chapter II), Regulation EU 2017/745

Basic UDI-DI: 8401934EC213L6

Common Specification: N/A

We herewith declare that the above mentioned products are in conformity with the Council Regulation 2017/745 for medical devices.

Compliance is applicable from the date listed below. All supporting documentation is retained at the premises of the manufacturer. This declaration is issued under the sole responsibility of ResMed Corp.

EC Certificate Number: G10 083904 0009 Rev.01

SRN: US-MF-000011805

Signed at: San Diego CA, USA on: 7th October 2025

Signed by:
Jason Gorman 07-Oct-2025 | 09:07 EDT
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Jason Gorman
Senior Director of Global Product Regulatory Affairs
ResMed Corp.

EC213.1
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