Declaration of Conformity

Manufacturer:	**Authorized Representative:**
ResMed Pty. Ltd. 1 Elizabeth Macarthur Drive Bella Vista NSW 2153 Australia	ResMed SAS Parc Technologique de Lyon 292 Allée Jacques Monod 69791 Saint Priest Cedex France

Product: RPSII- Air10 DC Cable

Intended Use:
The DC Cable is intended to connect battery pack to therapy device for discharging.

Classification: I according to Rule 1

EMDN: Z1203019080 Various Instruments for Anaesthesia and Pulmonary Ventilation Support - Hardware Accessories

Conformity Assessment Route: Annex II and III, Regulation EU 2017/745

Basic UDI-DI: 619498EC2086J

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 Pty. Ltd.

SRN: AU-MF-000011753

Signed at Sydney, Australia on: 12 March 2024

DocuSigned by:
Nicole Wilson
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Nicole Wilson
Person Responsible for Regulatory Compliance (PRRC)
ResMed Pty. Ltd.

EC 208a.4
First issued: 12 March 2024