



## Declaration of Conformity

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**Manufacturer:**

ResMed Corp.  
9001 Spectrum Center Blvd.  
San Diego CA 92123  
USA

**Authorized Representative:**

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

**Notified Body:**

TÜV SÜD Product Service  
GmbH  
Ridlerstraße 65  
80339 München  
Germany

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**Product:** AirView

**Intended Use:**

AirView is a web based solution for healthcare specialists intended to:

- assist in the diagnosis of sleep disordered breathing in adult patients through analysis of data recorded by an AirView compatible home sleep test device.
- transfer and display machine and therapeutic information that has been transmitted remotely from the patient's therapy device. It is intended to support the standard follow-up care of patients that have been prescribed a compatible ResMed therapy device. AirView also provides remote settings capabilities for non-life support devices only

**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:** 8401934EC137LF

**Common Specification:** N/A

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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

**SRN:** US-MF-000011805

Signed at San Diego CA, USA on: 14 November 2023

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Sheila Bruschi  
Director of Regulatory Affairs  
ResMed Corp.

**EC137.1**

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