ResMed

CE 0123

Declaration of Conformity

Manufacturer:	Authorised 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: 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 9

GMDN: 40582 Ventilator, software

Conformity Assessment Route: Annex II (excluding Section 4), 93/42/EEC

We herewith declare that the above mentioned products are in conformity with the Council Directive 93/42/EEC for medical devices including the MDD amendment 2007/47/EC.

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: G1 083904 0007

Signed at San Diego, USA on: 6 August 2020

Sheila Bruschi
Director of Regulatory Affairs
ResMed Corp.

EC137