RED Declaration of Conformity

Manufacturer:

ResMed Ltd
1 Elizabeth Macarthur Drive
Bella Vista
NSW 2153
Australia

European Representative:

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

Product Description:

Product Type:	AirMini
Trade Name(s):	ResMed
Model Number(s):	381XX
Software Version(s):	SW1.1.0.131
Accessories:	20W Power supply
Product Characteristics:	2400-2483.5 MHz; EIRP: <10dBm Channels: 79; Modulation: GFSK, π/4-DQPSK, 8-DPSK Channels: 40; Modulation: GFSK

Standards Applied:

Health *RED, Article 3.1a*	EN 62311:2008 EN 50566:2013/AC:2014 EN 62209-2:2010
Safety *RED, Article 3.1a*	EN 60950-1:2006+A11:2009+A1:2010+A12:2011+A2:2013
EMC *RED, Article 3.1b*	EN 301 489-1 V2.1.1 EN 301 489-17 V3.1.1
Radio Spectrum *RED, Article 3.2*	EN 300 328 V2.1.1

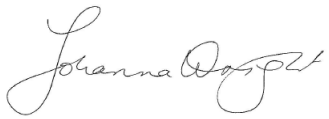
We herewith declare that the product meets national law of the essential requirements. The object of the declaration described above is in conformity with the relevant Union harmonization Legislation: Directive 2014/53/EU (RED).

All the reports of the applied standards have the positive opinion of Notified Body Cetecom with Notified Body number 0682 who issued the EU-type examination certificate.

Note: Any labelling of the product and printed material showing CE 0123, relates the Council Directive 93/42/EEC including the Medical Device Directive 2007/47/EC amendment.

This declaration is issued under the sole responsibility of the ResMed Ltd.

Signed at Sydney, Australia on: 25-Jul-18



Johanna Wright
Director of Regulatory Affairs
ResMed Ltd.