# RED Declaration of Conformity


**Manufacturer:**

ResMed Ltd
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Bella Vista
NSW 2153
Australia

**European Representative:**

ResMed SAS
Parc Technologique de Lyon
292 Allée Jacques Monod
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## *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<br>Channels: 79; Modulation: GFSK, π/4-DQPSK, 8-DPSK<br>Channels: 40; Modulation: GFSK |

## *Standards Applied:*

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

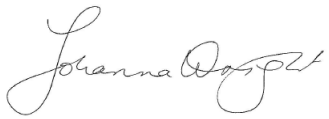
We herewith declare that the product meets national law of the essential requirements. The object of the declaration described above is in conformity with the relevant Union harmonization Legislation: Directive 2014/53/EU (RED).

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.