EC Declaration of Conformity

Manufacturer: ResMed Sensor Technologies Limited
NexusUCD Building,
Belfield Office Park,
Clonskeagh, Dublin 4. Ireland.

Device: **S+ by ResMed**
Model: **22111**
Description: **Sleep Monitor**

We ResMed Sensor Technologies Ltd declare that under our own responsibility our product **S+ by ResMed (Model No.: 22111**) to which this declaration relates is in conformity with the essential requirements and other relevant requirements of the

- R&TTE directive (1999/5/EC)
- The Low Voltage Directive (2006/95 /EC)
- The EMC Directive (2004/108/ EC)
- ROHS2 Directive (2011/65/EU).

The product is in conformity with the following standards and/or other normative documents:

Health (1999/5/EC Art 3.1(a)):	EN 62311:2008
Safety (1999/5/EC Art 3.1(a)):	EN 60950-1:2006+A11:2009+A1:2010+A12:2011+A2:2013
EMC (1999/5/EC Art 3.1(b)):	EN 301 489-1 V1.9.2,
	EN 301 489-3 V1.6.1,
	EN 301 489-17 V2.2.1
Spectrum (1999/5/EC Art 3.2):	EN 300 328 V1.9.1,
	EN 300 440-2 V1.4.1

The technical documentation relevant to the above equipment will be held at the address given above.

Signature _____ **Date March 9th, 2016**

Britta Jaffre
European Regulatory Affairs and Quality Operations Director
ResMed