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
69791 Saint Priest Cedex
France

Product Description:

Product Type:	Lumis VPAP S and ST Series (2G)
Trade Name(s):	ResMed
Model Number(s):	283xx
Software Version(s):	Product: SX584-xxxx, Cellular Module: SX558-xxxx
Accessories:	N/A
Product Characteristics:	GSM 900 MHz, 33dBm rated GSM 1800 MHz, 30dBm rated

Standards Applied:

Health *RED, Article 3.1a*	EN 62209-2:2010 EN 62311:2008
Safety *RED, Article 3.1a*	EN 60950-1:2006+A11:2009+A1:2010+A12:2011+A2:2013
EMC *RED, Article 3.1b*	Draft EN 301 489-1 V2.2.0 Draft EN 301 489-52 V1.1
Radio Spectrum *RED, Article 3.2*	EN 301 511 V9.0.2

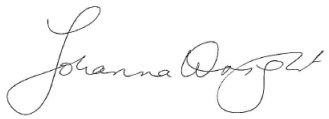
We herewith declare that the product meets national law of the essential requirements. The object of the declaration described above is in conformity with the relevant Union harmonization Legislation: Directive 2014/53/EU (RED).

All the reports of the applied standards have the positive opinion of Notified Body Telefication B.V. with Notified Body number 0560 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.

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