



# RED Declaration of Conformity

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**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

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**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

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**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

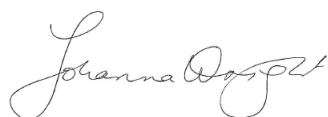
We herewith declare that the product meets national law of the essential requirements. The object of the declaration described above is in conformity with the relevant Union harmonization Legislation: Directive 2014/53/EU (RED).

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



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Johanna Wright  
Director of Regulatory Affairs  
ResMed Ltd.

**EC161**

First issue: 13-Jun-17