ResMed

Date: September 17, 2021

Re: Declaration about validity of test results from testing/certification of the integrated module

We,

ResMed Pty Ltd,
1 Elizabeth Macarthur Dr,
Bella Vista NSW 2153,
Australia

Related to product:

Type of Equipment:	Continuous Positive Airway Pressure (CPAP) Device
Brand name:	ResMed
Model name:	39485, 39486, 39487, 39488, 39489, 39490
Commercial name:	AirSense 11
FCC ID:	2ACHL-AIR11M1
IC:	9103A-AIR11M1

To whom it may concern,

We declare that the EXS62-W module has not been modified and has been integrated following module's manufacturer instructions for the installation such as input voltages considering extreme voltages, driver software, environmental conditions, etc...

Then, the results from conducted test reports according to FCC and ISED standards supplied by THALES DIS AIS Deutschland GmbH of module EXS62-W with *FCC ID QIPEXS62-W* and *IC 7830A-EXS62W* remain applicable, valid and representative of the module and are representative under the new conditions for the certification of the host device, AirSense 11

Test report Number	Test Standards	Date of Issue
191019009RFM-2R1 (LTE Report – Cat-M1)	FCC 47 CFR Part 22, FCC 47 CFR Part 24 FCC 47 CFR Part 27, FCC 47 CFR Part 90 RSS-130 Issue 2, RSS-132 Issue 3, RSS-133 Issue 6, RSS-139 Issue 3, RSS-Gen Issue 5	August 30, 2021

Sincerely,

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