Date: June 22, 2022

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:	37158, 37159, 37160, 37161, 37162, 37163, 37164, 37165
Commercial name:	AirSense 10, AirCurve 10
FCC ID:	2ACHL-AIR104GU
IC:	9103A-AIR104GU

To whom it may concern,

We declare that the u-blox LARA R6001D 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 u-blox of module LARA R6001D with *FCC ID XPYUBX21BE01* and *IC 8595A-UBX21BE01* remain applicable, valid and representative of the module and are representative under the new conditions for the certification of the host device.

Sincerely,

Christopher Jenkins

By:	Christopher Jenkins
Title:	Associate Manager – Systems Engineering
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