Date: January 25, 2023

Re: Declaration about validity of test results from testing/certification of the previous model

Addressed to: Federal Communications Commision
Equipment Authorization Branch
7435 Oakland Mills Rd.
Columbia MD 21046, USA

Issued by: 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:	39523,39524,39525,39526,39527,39528
Commercial name:	AirSense 11 AutoSet USA, AirSense 11 CPAP USA, AirSense 11 Elite USA, AirSense 11 AutoSet CAN, AirSense 11 CPAP CAN, AirSense 11 Elite CAN
FCC ID:	2ACHL-AIR11M1U

To whom it may concern,

We declare that the model 39485 with HW version: R390-7654 and the model 39485 with HW version: R390-7667 support the same module BLE as it can be confirmed with the electrical diagram and list of materials sent with along this declaration.

Then, the results from conducted test according to FCC and ISED standards remain applicable, valid and representative for the certification of the host device, AirSense 11

Test report Number	Test Standards	Date of Issue
72943RRF.005A1	USA FCC Part 15.247, 15.209 CANADA RSS-247, RSS-Gen	2022-12-23

Sincerely,

By: Christopher Jenkins
Title: Manager – Systems Engineering
Company: ResMed Pty Ltd
Telephone: +61 2 8884 1517
e-mail: Christopher.Jenkins@resmed.com.au