



**Date:** February 12, 2024

**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:        | 28541, 28542, 28405                               |
| Commercial name:   | AirCurve 10 ST-A                                  |
| 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,

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