



## Troubleshooting Guide for AIRVO

This Troubleshooting Guide is intended for technical users, including clinical/biomedical engineers and technical personnel, of the AIRVO™ 2 humidifier. It applies to all AIRVO 2 humidifiers from lot numbers **140910** and above.

If this troubleshooting guide does not resolve your issue, please contact your local Fisher & Paykel Healthcare representative.

### D.1 AIRVO does not turn on

- A. Press and hold the ON/OFF button for at least 2 seconds.
- B. Is the AIRVO 2 plugged into mains power?
- C. Is the power cord securely inserted into the back of the AIRVO 2?
- D. Is the power cord damaged?
  - If yes, replace the damaged cord. See **Section 5.1** for a **900PT410xx** replacement power cord.
  - If no, try using another power cord.
- E. Connect the AIRVO 2 into another power outlet.
- F. Connect a different electrical device into the same power outlet. Turn on the device to confirm that the power outlet is working.
- G. The AIRVO 2 may be 'on' with a broken display.  
Turn the AIRVO 2 on without the heated breathing tube and check that the audible alarm activates.

### D.2 Power out (black screen)

The auditory alarm will sound for at least 120 seconds.

*The most likely cause is a dislodged or disconnected power cord.*

- A. Please follow the instructions in **Section D.1**.  
Note: Press "audio pause" button to permanently silence the alarm (  ).  
The device will not automatically restart.

### D.3 "Check tube"<sup>Fig. 1</sup> or "E38"

- A. Is the heated breathing tube attached correctly?
  - Even if it appears to be, unplug and reconnect the heated breathing tube.
- B. Is the heated breathing tube visibly damaged?
  - Check the electrical pins and the tube itself.
- C. Try using a new heated breathing tube.



Figure 1



## D.4 “Check for blockages”<sup>Fig. 2</sup> or “E121”

### D.4.1 WATER CHAMBER AND NON-RETURN VALVE

- A. Have the silicone flaps of the non-return valve, found inside the left-hand chamber port, been displaced<sup>Fig. 3</sup>?
- If yes, return them to the correct position using a non-sharp tool, such as a pair of non-sharp tweezers<sup>Fig. 5</sup>.

Note: If the Non-return valve is damaged or missing, replace with part **900PT911**. Upon replacement, ensure the spine is sitting vertically<sup>Fig. 5</sup>. If placed horizontally, this may cause the bottom flap to open due to gravity<sup>Fig. 4</sup>. This may cause both “Check for leaks” and “Check for blockages” warnings.

- B. Is the MR290 water chamber overfilled above the black line?
- If yes, replace with a new water chamber. Contact your local Fisher & Paykel Healthcare representative about the faulty chamber.

### D.4.2 HEATED BREATHING TUBE

- A. Is the heated breathing tube visibly blocked or kinked<sup>Fig. 6</sup>?

### D.4.3 PATIENT INTERFACE AND AIRVO MODE

- A. Is the patient interface visibly blocked or kinked?
- B. Should the unit be in Junior mode<sup>Fig. 7</sup>?
- If the AIRVO is in Default mode and the 900PT531 Junior tube is used with the OPT316 and OPT318 cannula interfaces it may generate a “Check for blockages” alarm.
- See **Appendix E** for identification of the Default and Junior Tube and Chamber Kits, according to their labels.
- C. Are you using an unsuitable cannula?
- The OPT312 and OPT314 cannot be used with the AIRVO 2.
- See the User Manual for information regarding patient interfaces.

### D.4.4 AIR FILTER

- A. Is the air filter significantly discolored/dirty?
- Replace with part **900PT913**.



Note: A prompt<sup>Fig. 8</sup> for filter change will occur at the start of the Disinfection Cycle once the AIRVO 2 has counted 1,000 hours of use. Choose ‘Now’ or ‘Later’<sup>Fig. 9</sup> by using the “up” or “down” buttons and press the “mode” button (▶) to confirm. Selecting ‘Now’ will zero the counter. Selecting “Later” will activate the prompt at the start of the next Disinfection Cycle.

- B. Is there a foreign object blocking the air filter or filter holder?

### D.4.5 CONDENSATION

Please see **Section D.12**.

### D.4.5 ALTITUDE

- A. The myAIRVO 2 is designed to operate at an altitude below 2,000 meters.



Figure 2

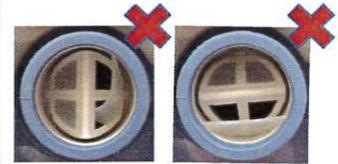


Figure 3

Figure 4



Figure 5

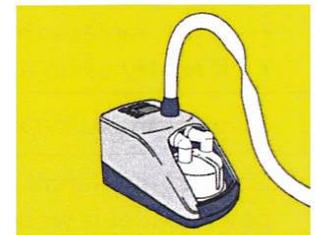


Figure 6

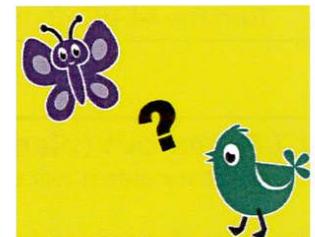


Figure 7



Figure 8



Figure 9



## D.5 “Check for leaks”<sup>Fig. 10</sup> or “E122”

The most likely cause is a missing water chamber or the existing chamber has not been pushed into place correctly.

### D.5.1 WATER CHAMBER

- A. Is the water chamber fitted correctly? Even if it appears to be:
- Remove the water chamber.
  - Push the chamber on firmly, until the finger guard “clicks” into place<sup>Fig. 11</sup>.
- ⚠ Warning: The heater-plate and base of the water chamber may be hot.

### D.5.2 HEATED BREATHING TUBE

- A. Is the heated breathing tube attached to the device correctly? Even if it appears to be:
- Disconnect the heated breathing tube.
  - Check that the black O-ring is in place<sup>Fig. 12</sup>. If the O-ring is damaged or missing, replace with part **900PT408**.
  - Reconnect the heated breathing tube.

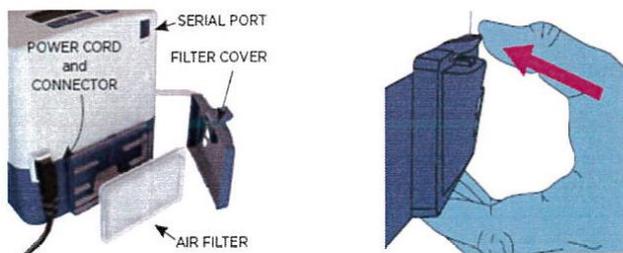
- B. Confirm that the heated breathing tube is not visibly damaged.

### D.5.3 PATIENT INTERFACE

- A. Is the patient interface correctly fitted to the heated breathing tube? Even if it appears to be, disconnect and reconnect the patient interface. It should make a “click” sound when it is connected properly.
- B. Should the unit be in Default (adult) mode?
- If the AIRVO is in Junior mode and the 900PT501 Default tube is used with the OPT842/44/46/70 or RT013 interfaces, it may generate a “Check for leaks” alarm.
- See **Appendix E** for identification of the Default and Junior Tube and Chamber Kits, according to their labels.

### D.5.4 AIR FILTER & FILTER COVER

- A. Is the air filter and filter cover (at the back of the device) correctly fitted, as per the User Manual?



Check for leaks

Figure 10

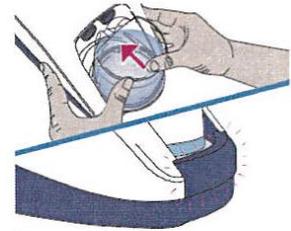


Figure 11

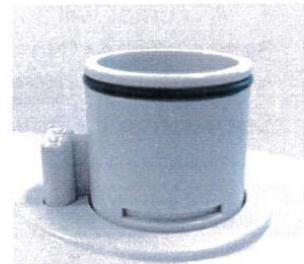


Figure 12



### D.6 “O<sub>2</sub> too low”<sup>Fig. 13</sup>

The measured oxygen level has fallen below the allowed limit.

- Adjust the level of oxygen from the oxygen source as necessary, i.e. increase the oxygen flow rate through the oxygen flow meter.
- Is the oxygen source (wall or cylinder flow meter) turned on?
- Is the oxygen source empty or faulty?
- Is the “AIRVO 2 oxygen inlet kit”<sup>Fig. 14</sup> installed correctly, as per the instructions included with part **900PT422** and confirmed that there are no kinks in the “AIRVO 2 oxygen inlet kit” oxygen tubing?
- Is the oxygen source tubing correctly and securely fitted to the AIRVO 2?
- Allow the device to sufficiently warm up; rapid changes in temperature can affect the sensor.
- Is the minimum oxygen limit set to 25%?
  - A prompt will appear with an option to change this lower limit to 21%. Select “Yes” or “No” by using the “Up” and “Down” buttons. Press the “mode” button (  ) to confirm selection<sup>Fig. 15</sup>.
 See **Section 2 - Advanced Settings** to change this lower oxygen limit.



Figure 13



Figure 14



Figure 15

### D.7 “O<sub>2</sub> too high”<sup>Fig. 16</sup>

The measured oxygen level has risen above the allowed limit.

- Adjust the level of oxygen from the oxygen source as necessary, i.e. decrease the oxygen flow rate through the oxygen flow meter.
- See **Section 2 - Advanced Settings** to change this lower oxygen limit.



Figure 16

### D.8 “Cannot reach target flow”<sup>Fig. 17</sup>

- Press the “mode” button (  ) to continue normal operation at a lower (maximum achievable) flow rate.
- Is the target flow setting too high for the patient interface?
  - Check the swing tag/User Manual for the appropriate flow range for each patient interface.
 Note: If the AIRVO 2 cannot reach the target flow setting, it will automatically select a maximum achievable flow rate and prompt the user to press the “mode” button (  ) to confirm.
- Follow steps in **Section D.4** – “Check for blockages”.
- Is the altitude above 2,000 m?  
The AIRVO 2 is designed to operate at an altitude below 2,000 meters.



Figure 17



<p><b>D.9 “Cannot reach target temperature”</b> Fig. 18</p> <p><i>The most likely cause is operating the AIRVO 2 at a high flow rate in a cold room. Consider decreasing the target flow setting.</i></p>	 <p><b>Cannot reach target temperature</b></p> <p>Figure 18</p>
<p>A. Press “mode” button (  ) to continue. Note: The humidity level may be compromised.</p>	
<p>B. Is the ambient room temperature below 18 °C (64 °F)?</p> <ul style="list-style-type: none"> <li>• If yes, proactive management of condensation may be required. See <b>Section D.12</b> on prevention and management of condensation.</li> </ul>	

<p><b>D.10 “Check water”</b> Fig. 19</p>	 <p><b>Check water</b></p> <p>Figure 19</p>
<p>A. Is the water bag empty? If yes, refill or replace the water bag and press the “mode” button (  ) to reset the alarm.</p>	
<p>B. Is the water chamber empty?</p> <ul style="list-style-type: none"> <li>• If yes, replace the water chamber as it may be damaged.  Warning: The heater-plate and base of the water chamber may be hot.</li> </ul>	
<p>C. Is there a kink in the fluid line, preventing water from flowing into the chamber?</p>	
<p>D. Open the vent cap near the water bag spike. This allows the pressure to equalize, letting the water flow into the water chamber.</p>	

<p><b>D.11 “Check operating conditions”</b> Fig. 20</p> <p><i>This alarm may be caused by a sudden change in ambient room temperature, e.g. storing the unit in a cold place, then using it in a warm place.</i></p>	 <p><b>Check operating conditions</b></p> <p>Figure 20</p>
<p>A. Is the ambient room temperature less than 10 °C (50 °F) or greater than 30 °C (86 °F)?</p>	
<p>B. Leave the unit running for 30 minutes. Switch the unit off, then restart.</p>	



## D.12 Condensation

### D.12.1 PREVENTION OF EXCESSIVE CONDENSATION

- A. Is the AIRVO 2 being used in ambient conditions between 18 - 28 °C (64 - 82 °F)?
- If the room is less than 18 °C (64 °F), condensation is more likely to occur.
- B. Is there a local source of cooling acting on the heated breathing tube?
- A fan to cool the patient,
  - An air-conditioning unit, vent or an open window?
  - Are you able to remove or minimize these sources of cooling, e.g. redirect the fan, cooling the patient, away from the heated breathing tube?

### D.12.2 CONDENSATION MANAGEMENT

- A. Implement a system to check the heated breathing tube for condensate regularly.
- B. Is the AIRVO 2 placed below head height<sup>Fig. 21</sup>?
- This will allow condensate to drain towards the water chamber, away from the patient.
- C. If condensation is present, drain it back into the water chamber<sup>Fig. 22</sup>:
- Disconnect the patient interface from the heated breathing tube.
  - Drain the tube by lifting the patient end of the tube, allowing the condensate to run into the water chamber.
  - At higher target flow rates, it may be necessary to first reduce the target flow rate to 30 L/min or below, to ensure the condensate drains into the water chamber.
- D. If condensate persists, consider turning the target temperature down.
- A lower target temperature will decrease the humidity output of the AIRVO 2, decreasing the level of condensation.
- Note: The temperature and humidity level delivered to the patient will also be reduced.

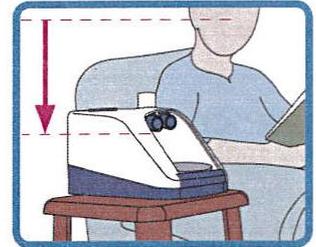


Figure 21

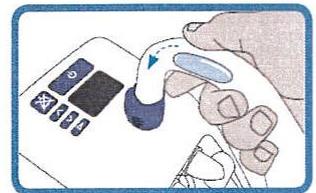


Figure 22