ResMed

AirSense™ 11

AUTOSET

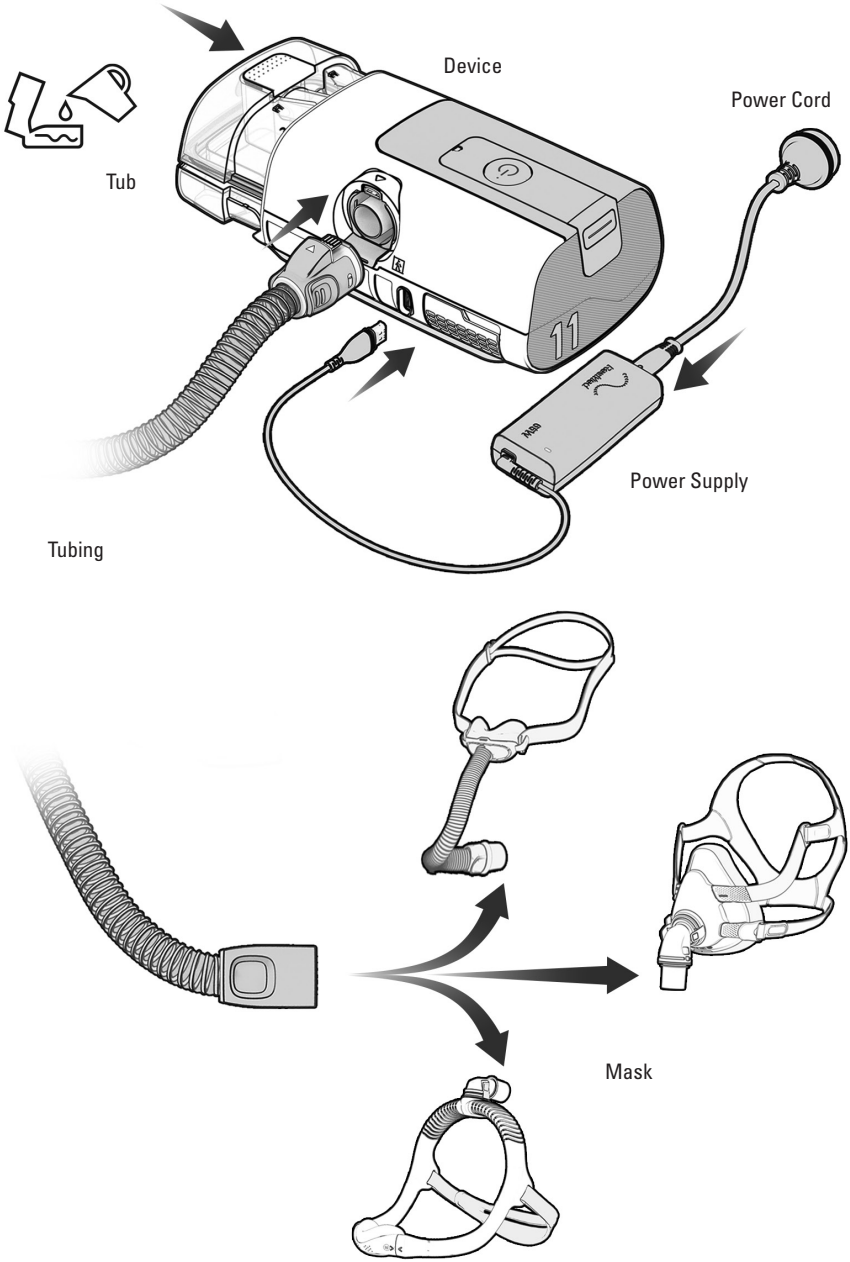
CPAP

ELITE

User guide

English

Setting Up



Contents

Welcome

The AirSense™ 11 AutoSet™ (including AutoSet for Her) device is ResMed's premium auto-adjusting pressure device. The AirSense 11 Elite and the AirSense 11 CPAP are ResMed's Continuous Positive Airway Pressure (CPAP) devices.

⚠ WARNING

Read this entire guide before using the device.

⚠ CAUTION

In the US, Federal law restricts this device to sale by or on the order of a physician.

Indications for use

AirSense 11 AutoSet (including AutoSet for Her)

The AirSense 11 self-adjusting system is indicated for the treatment of obstructive sleep apnea (OSA) in patients weighing more than 66 lb (30 kg), including female patients with mild to moderate OSA in AutoSet for Her mode. The AirSense 11 self-adjusting system is intended for home and hospital use.

AirSense 11 CPAP (including Elite)

The AirSense 11 CPAP system is indicated for the treatment of obstructive sleep apnea (OSA) in patients weighing more than 66 lb (30 kg). The AirSense 11 CPAP system is intended for home and hospital use.

Contraindications

Positive airway pressure therapy may be contraindicated in some patients with the following pre-existing conditions:

- severe bullous lung disease
- pneumothorax or pneumomediastinum
- pathologically low blood pressure, particularly if associated with intravascular volume depletion
- dehydration
- cerebrospinal fluid leak, recent cranial surgery, or trauma.

Adverse effects

You should report unusual chest pain, severe headache, or increased breathlessness to your prescribing physician. An acute upper respiratory tract infection may require temporary discontinuation of treatment.

The following side effects may arise during the course of therapy with the device:

- drying of the nose, mouth, or throat
- nosebleed
- bloating
- ear or sinus discomfort
- eye irritation
- skin rashes.

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At a glance

⚠ WARNING

Use only recommended ResMed masks and accessories or other vented masks as recommended by the prescribing doctor with this device. Using these components allows normal breathing and prevents potential asphyxiation.

The AirSense 11 system includes the following:

- Device
- HumidAir™ 11 Standard tub
- ClimateLineAir™ 11 heated tubing or SlimLine™ tubing
- 65W AC adaptor
- Travel bag
- SD card (not available in all devices).

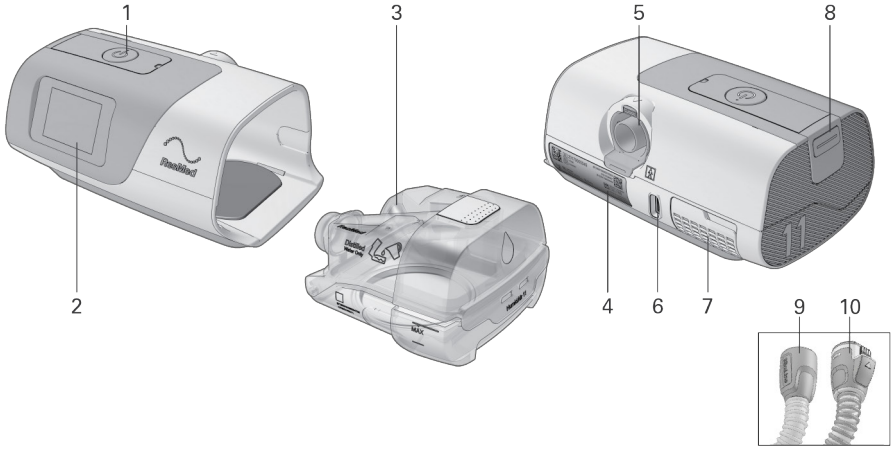
Contact your care provider for a range of accessories available for use with the device including:

- Air tubing (ClimateLineAir 11 and SlimLine)
- HumidAir 11 Standard tub
- HumidAir 11 Cleanable tub (can be disinfected)
- End cap which allows use without the humidifier
- Air11™ Filter - standard
- Air11 Filter - hypoallergenic
- SD card

Notes:

- The AirSense 11 device is compatible with ResMed masks. For a full list, see the Mask/Device compatibility list on ResMed.com/downloads/devices.
- The HumidAir 11 Standard tub and HumidAir 11 Cleanable tub are the only water tubs used with the AirSense 11 device.
- The ClimateLineAir 11 is the only heated tubing that is compatible with the AirSense 11 device.

About your device



	Description	Purpose
1	Start Therapy/ Standby button	Press to start/stop therapy. The LED indicator is green during standby mode, and white during therapy, **Test Drive,** and **Mask Fit** functions.
2	Display touch screen	Navigates between functions and displays information on the operating status of the device.
3	HumidAir 11 tub	Water tub that provides heated humidification.
4	Device label	Contains information relevant to the device.
5	Outlet connector	Connects the air tubing
6	Power inlet	Connects the power cord
7	Air inlet filter cover	Contains the air filter
8	SD card cover	Removable cover that protects the SD card slot. The LED indicator is blue when data is written to the SD card.
9	SlimLine tubing	Non-heated air tubing
10	ClimateLineAir 11 tubing	Heated air tubing

Notes:

- If the Start therapy/ Standby button has a flashing white light, a system error has occurred. Refer to the Troubleshooting section for more information.
- Use this device only as directed by your physician or healthcare provider.

Setting up your device

⚠ WARNING

Do not use any additives in the humidifier tub (eg, scented oils or perfumes). These may reduce humidification output and/or cause deterioration of the tub materials.

⚠ CAUTION

Use only ResMed parts (eg air inlet filter, power supplies), masks and accessories with the machine. Non ResMed parts may reduce the effectiveness of the treatment, may result in excess carbon dioxide rebreathing and/or damage the machine. For compatibility information, refer to ResMed.com for more information.

When using the humidifier tub:

- Always place the device on a level surface, lower than your head, to prevent the mask and air tubing from filling with water.
- Do not overfill the humidifier tub as water may enter the device and air tubing.
- Do not fill the humidifier tub with hot water as this could lead to excessive air temperature at the mask. Ensure the water is cooled to room temperature before filling the humidifier tub.
- Do not place the device on its side while the humidifier is attached as water might get into the device and reduce motor life.

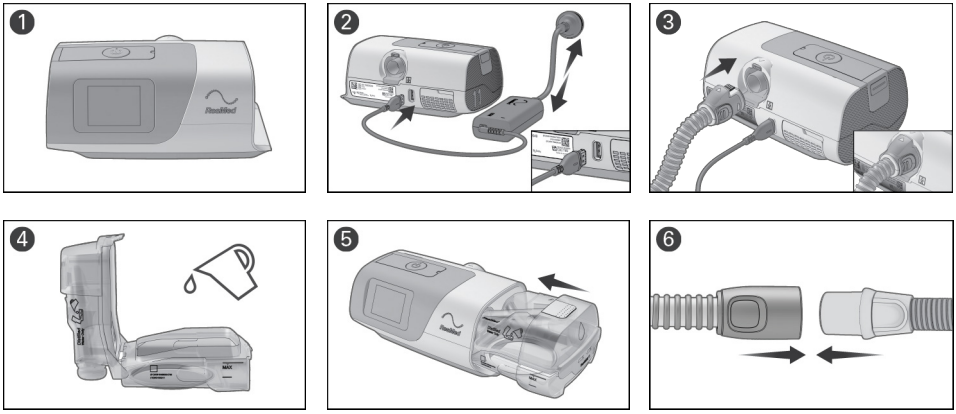
When setting up the AirSense system:

- Do not place the power supply where it can be bumped, stepped on, or where someone is likely to trip over the power cord.
- Do not block the air tubing and/or air inlet of the device while in operation as this could lead to overheating of the device.
- Keep the area around the device dry, clean and clear of anything (eg, clothes or bedding) that could block the air inlet or cover the power supply unit.
- Ensure the system is correctly setup. Incorrect system setup may result in incorrect mask pressure reading.

When using a mask:

- Use only vented masks recommended by ResMed or by the prescribing doctor with this device.
- Fitting the mask without the device blowing air can result in rebreathing of exhaled air.
- Make sure that the mask vent holes are kept clear and unblocked to maintain the flow of fresh air into the mask.

To set up the device:



1. Place the device on a stable level surface.
2. Connect the power cord into the power inlet at the rear of the device. Connect one end of the power cord into the AC adaptor and the other end into the power outlet.
3. Connect the air tubing firmly to the outlet connector at the rear of the device.
4. Open the humidifier tub and fill it with distilled water under room temperature up to the maximum water level mark. The humidifier tub must be removed from the device before adding water. The humidifier tub has a maximum capacity of 380 mL.
5. Close the humidifier tub and insert it into the side of the device.
6. Connect the free end of the air tubing firmly onto the assembled mask.
   See the mask user guide for detailed information.

Recommended masks for use with this device are listed on Resmed.com

Notes:

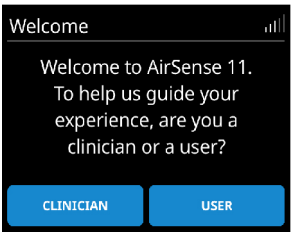
- Do not insert any USB cable into the AirSense 11 device or attempt to plug the AC adaptor into a USB device. This may cause damage to the AirSense 11 device or USB device.
- The electrical connector end of the heated air tubing is only compatible with the air outlet at the device end and should not be fitted to the mask.
- Do not use electrically conductive or anti-static air tubing.

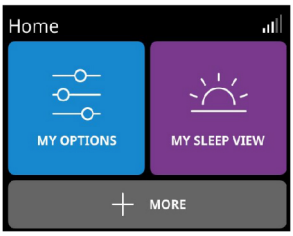
Navigating the touch screen

The AirSense 11 device operates via a display touch screen, which allows you to access, view and change therapy and device settings. You can also track your sleep health progress.

The status bar at the top of the screen may display icons at different times and may include:

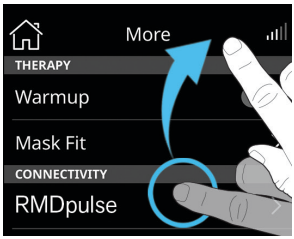
Icon	Description	Purpose
	Home Screen	Return to the Home screen at any time.
	Humidifier fault	Detects fault in the humidifier. Therapy will run without heating.
	Humidifier warming	Water in the humidifier tub is pre-heating.
	Humidifier cooling	Water in the humidifier tub is cooling.
	Bluetooth® connected	Device is successfully connected via Bluetooth.
	Cellular signal strength	Indicates the strength of cellular connectivity
	No cellular connection	Cellular coverage is not available
	Airplane mode	Device is in airplane mode

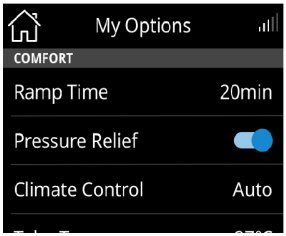




1. From the **Welcome** screen, tap **USER**
2. From the **Home** screen, you can access the following menus:
   - **MY OPTIONS**: View and adjust therapy settings (eg, Adjust Ramp time)
   - **MY SLEEP VIEW**: Track sleep health (check the number of hours used last night or mask status)
   - **MORE:** Access additional features such as Run Mask Fit or switch to Airplane mode.

Using the touch screen:





There are two actions to navigate through the touch screen:

Flick: Flick up or down the screen to view menu options. A flick action will move the menu one full screen.

Tap: Select a parameter setting to update. For other parameters (eg Pressure Relief, Airplane mode), tap the parameter to turn it on  or tap to turn it off  .

Personalizing your settings

The device has been set up for your needs by your care provider, but you may want to make adjustments to make your therapy more comfortable.

1. Tap **MY OPTIONS** from the **Home** screen.
2. Tap the parameter you wish to change.
3. Tap the preferred setting.
   Tap **OK** to confirm the change or **CANCEL** to go back to the previous screen.

Additional features

There are some other features on your device which you can personalize.

Menu	Function	Description
MY OPTIONS	Ramp Time	Period during which the pressure increases from a low start pressure to the prescribed treatment pressure. Ramp Time can be set to Off, 5 to 45 minutes (in 5-minute increments), or Auto.
	Pressure Relief*	When EPR is enabled, you may find it easier to breathe out. This setting can help you get used to therapy.
	SmartStart*	When SmartStart is enabled, therapy starts automatically when you breathe into your mask.
	SmartStop*	When SmartStop is enabled, therapy stops automatically after a few seconds when you remove your mask.
MORE	Mask Fit	This function helps you assess and identify possible air leaks around your mask.

*Features enabled by your care provider.

Note: Not all functions are available in all regions. Functions vary based on therapy mode.

Connecting your AirSense 11 device and smart device

RMDpulse is a smartphone app that guides you through the setup process. This includes device setup videos, mask fitting videos, trying therapy using the Test Drive feature, and tracking your sleep health progress. The app is not required to operate the AirSense 11 device.

Before pairing the AirSense 11 device to a smartphone, ensure the app's latest version is installed on the smartphone. If not, download the app from the App Store® or on Google® Play. Pair the AirSense 11 device to your phone. To set up the app, go to the MORE menu.

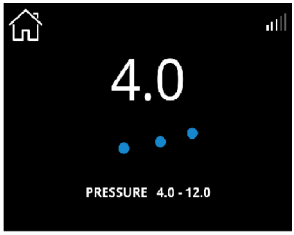
1. Ensure your AirSense 11 device is set up correctly and plugged into a power source.
2. Launch the RMDpulse app. Tap **Continue.**
3. Follow the prompts on the RMDpulse app to complete the BT connection.
   AirSense 11 is now connected to the app. The Bluetooth connection symbol appears on the status bar to confirm the connection between the AirSense 11 device and the smartphone.
4. Tap **Done**.

Starting/Stopping therapy

⚠ WARNING

The machine is not intended to be operated by persons (including children) with reduced physical, sensory or mental capabilities without adequate supervision by a person responsible for the patient's safety.

To start therapy:



1. Fit your mask
2. Press the Start therapy/Standby button or breathe normally if SmartStart is enabled.

Therapy will begin and the Treatment screen is displayed. A dynamic pulse wave will appear during therapy.

Notes:

- The screen will fade and then go black automatically after a short period of time. Tap the screen to turn it back on.
- If power is interrupted during therapy, the device will automatically restart therapy when power is restored.
- The device has a light sensor that adjusts the screen brightness based on the light in the room.

To stop therapy:

1. Remove your mask.
2. Press the Start/Standby button or wait until the device stops if SmartStop is enabled.

About the heated tubing

The ClimateLineAir 11 is a heated breathing tube that delivers air to a compatible mask. When used with the device humidifier tub, ClimateLineAir 11 heated air tubing allows you to use the Climate Control feature.

Note: Not all types of air tubing are available in all regions.

Climate Control

Climate Control is designed to make therapy more comfortable by enabling constant temperature and maintaining humidity.

This feature:

- delivers comfortable humidity level and temperature during therapy
- maintains the set temperature and relative humidity during sleep to prevent dryness in the nose and mouth
- can be set to either Auto or Manual
- is only available when both the ClimateLineAir 11 and HumidAir tub are attached.

Climate Control - Auto setting

Auto is the recommended and default setting. It is designed to make therapy as easy as possible so there is no need to change the temperature or humidity settings.

- Sets the tube temperature to Auto (80ºF/27ºC). If the air in the mask is too warm or too cold, you can adjust the tube temperature to anywhere from 60 to 86ºF (16 to 30ºC) or turn it off completely
- Adjusts the humidifier output to maintain a constant, comfortable humidity level of 85% relative humidity
- Protects against rainout (water droplets in the heated air tubing and mask).

Climate Control - Manual setting

Manual is designed to offer more flexibility and control over settings and offers the following:

- Temperature and humidity can be adjusted to find the most comfortable setting
- Temperature and humidity level can be set independently
- Rainout protection is not guaranteed. If rainout does occur, first try increasing the tube temperature
- If the air temperature becomes too warm and rainout continues, try decreasing the humidity.

Note: If Climate Control is set to Manual, the Auto Tube Temperature setting is not available.

Humidity Level

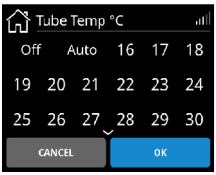
The humidifier moistens the air and is designed to make therapy more comfortable.

- If you are getting a dry nose or mouth, turn up the humidity
- If you are getting any moisture in your mask, turn down the humidity.
- You can set the Humidity Level to Off or between 1 and 8, where 1 is the lowest humidity setting, and 8 is the highest humidity setting.

To update the setting for Tube Temperature, Climate Control, or Humidity Level, tap MY OPTIONS from the Home screen, go down the list of options, and select the setting.

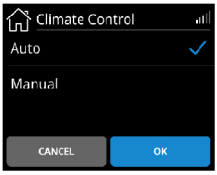
Note: Tube Temp Auto setting is only relevant when using the Climate Control Auto setting. If Climate Control is set to Manual, Auto set temperature is not a valid selection.

Tube Temperature



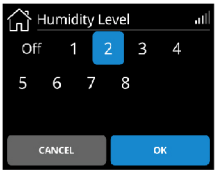
1. Tap **Tube Temp**.
2. Tap the preferred setting.
3. Tap **OK** to save the change.

Climate Control



1. Tap **Climate Control.**
2. Tap **Manual**.
3. Tap **OK** to save the change.

Humidity Level



1. Tap **Humidity Level.**
2. Tap the preferred setting.
3. Tap **OK** to save the change.

Note: The temperature and humidity settings are not measured values.

Therapy data

The AirSense 11 device records your therapy data for viewing and adjusting by your care provider if required. The data is transferred to your care provider in the following methods:

Wireless

The device is equipped with cellular communication that allows your sleep therapy data to be wirelessly transmitted to your care provider. It also allows for your therapy settings to be updated.

Transfer of data will occur after therapy has stopped. Leave your device connected to the power outlet at all times and make sure it is not in Airplane Mode. Data will only be transferred if a wireless connection is available.

Please be aware that within the wireless network, the availability and quality of the network may be affected by terrain, buildings, and the weather. Wireless communication depends on network availability. Coverage is not available everywhere and varies by service.

Notes:

- Therapy data might not be transmitted if you use it outside of the country or region of purchase.
- Devices with cellular communication might not be available in all regions.

SD card

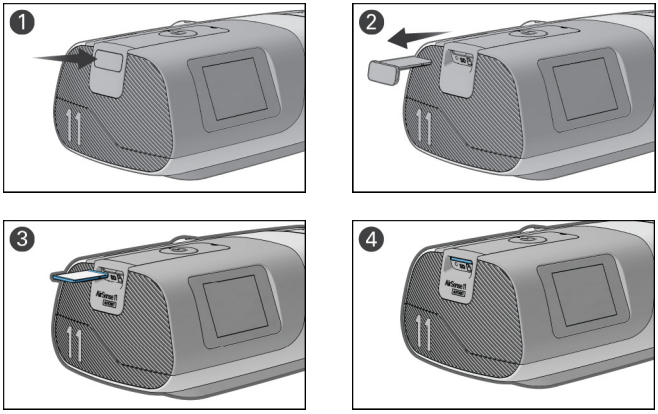
Your sleep therapy data may be transferred to your care provider via SD Card (if provided). Your care provider may ask you to send the SD card by mail or to bring it in. Only remove the SD card when instructed by your care provider.

To use the SD card to record your sleep data, remove the SD card cover.

Do not remove the SD card from the device when the SD light is flashing, because data is being written to the card.

Note: The SD card should not be used for any other purpose as it may corrupt therapy data stored on the card.

To remove the SD card cover and insert SD card:



1. Push the SD card cover.
2. Remove the SD card cover and keep the SD card cover in a safe place.
3. Insert the SD card.
4. Push in the SD card until it clicks in place.

To remove the SD card:

1. Push in the SD card to release it.
2. Place the SD card in the protective folder and follow your care provider's instructions.

For more information on the SD card, refer to the SD card protective folder provided with your device.

Cleaning and caring for the device

⚠ WARNING

- Beware of electrocution:
  - Do not immerse the device, AC Adaptor or power cord in water.
  - Do not connect to power while the device is wet. Make sure that all parts are dry before plugging it in.
  - If liquids are spilled into or onto the device, unplug the device and let the parts dry.
- Always unplug the device before cleaning and ensure that all parts are dry before plugging it back in.
- Do not perform any maintenance tasks (eg, cleaning, changing the air filter) while the device is in operation.
- Clean the device and its components according to the schedules shown in this guide, to maintain the quality of the device and to prevent the growth of germs that can adversely affect your health.
- Regularly inspect power cords, cables, and power supply for damage or signs of wear. Discontinue use and replace if damaged.
- Do not open or modify the device. There are no user serviceable parts inside. Repairs and servicing should only be performed by an authorized ResMed service agent.

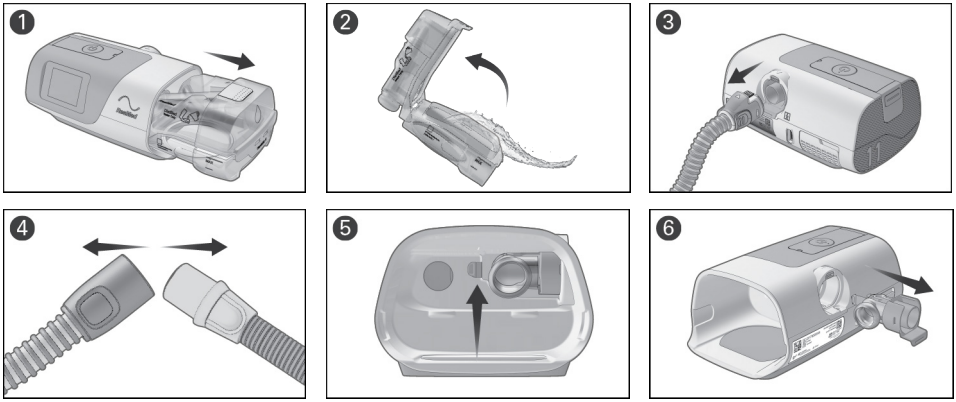
⚠ CAUTION

- Do not use bleach, chlorine, alcohol, or aromatic-based solutions, moisturizing or antibacterial soaps or scented oils to clean the device, the humidifier tub or air tubing. These solutions may cause damage or affect the humidifier performance and reduce the life of the products. Exposure to smoke, including cigarette, cigar or pipe smoke, as well as ozone or other gases, may damage the device. Damage caused by any of the foregoing, will not be covered by ResMed's limited warranty.
- Leave the humidifier tub to cool for ten minutes before handling to allow the water to cool and to make sure that the humidifier tub is not too hot to touch.

The following sections will help you with:

- Disassembling
- Cleaning
- Checking
- Reassembling.

Disassembling



1. Hold the humidifier tub at the top and bottom, press it gently and pull it away from the device.
   Note: Take care when handling the humidifier tub as the humidifier tub may be hot. Allow 10 minutes for the heater plate and any excess water to cool.
2. Open the humidifier tub and discard any remaining water.
3. Pinch the cuff of the air tubing, and gently pull it away from the device.
4. Hold both the cuff of the air tubing and the swivel of the mask, then gently pull apart.
5. Locate the outlet connector on the inside of the device and release it by pressing the clip firmly.
6. Remove the outlet connector by pulling it out through the outlet connector socket at the rear of the device.

Cleaning

The following instructions are for home cleaning. Instructions for reprocessing devices intended for multi-patient re-use can be found in the clinical guide.

You should clean the device, humidifier tub, air tubing, and outlet connector as described.

Daily:

1. Empty the humidifier tub and wipe it thoroughly with a clean disposable cloth. Allow it to dry out of direct sunlight.
2. Refill the humidifier tub with distilled water.

Weekly:

1. Wash the humidifier tub, air tubing, and outlet connector in warm water using a household dishwashing liquid. They should not be washed in temperatures higher than 149°F (65°C).
2. Rinse each component thoroughly in water.
3. Allow to dry out of direct sunlight or heat.
4. Wipe the exterior of the device with a dry cloth.

Notes:

- The humidifier tub may be washed in a dishwasher on the delicate cycle (top shelf only).
- Do not wash the air tubing in a dishwasher or washing machine.
- The air filter is not washable or reusable.

For cleaning your mask, refer to the mask user guide for detailed instructions.

Checking

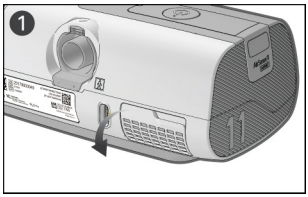
⚠ WARNING

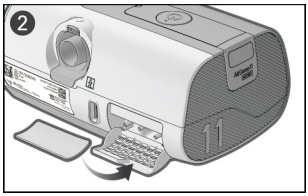
- Discontinue use and contact your care provider or ResMed Service Center if any of the following occur:
  - device does not perform as usual
  - device is making unusual sounds
  - device is damaged
- If any visible deterioration of a system component is apparent (cracking, discoloration, tears etc.), the component should be discarded and replaced.
- If used, regularly check the bacterial/viral filter for signs of moisture or other contaminants, particularly during nebulization or humidification. Failure to do so could result in increased breathing resistance.

Regularly check the humidifier tub, air tubing, and air filter for any damage.

1. Check the humidifier tub:
   - Replace it if it is leaking or has become cracked, cloudy, or pitted.
   - Replace it if the seal is cracked or torn.
   - Remove any white powder deposits using a solution of one-part household vinegar to 10 parts water. Rinse with clean water.
2. Check the air tubing and replace it if there are any holes, tears or cracks.
3. Check the air filter and replace it at least every six months. Replace it more often if there are any holes or blockages by dirt or dust.

Replacing the air filter





1. Open the air filter cover and remove the old air filter. The air filter is not washable or reusable.
2. Place a new air filter onto the air filter cover and then close the cover. Make sure the air filter and air filter cover is fitted at all times to prevent water and dust from entering the device.

Reassembling

When the humidifier tub and air tubing are dry, you can reassemble the parts.

To reassemble the AirSense 11 system:

1. Hold the outlet connector with the seal pointing to the left and the clip pointing forward.
2. Make sure the outlet connector is correctly aligned and insert the outlet connector into the socket.
3. Check the outlet connector is inserted correctly.
4. Connect the air tubing firmly to the air outlet located on the rear of the device..
5. Open the humidifier tub and fill it with distilled water under room temperature up to the maximum water level mark.
6. Close the humidifier tub and insert it into the side of the device.
7. Connect the free end of the air tubing firmly onto the assembled mask.

Traveling

You can take your device with you wherever you go. Just keep the following in mind:

- Use the travel bag provided to prevent damage to the device.
- Empty the humidifier tub and pack it separately in the travel bag.
- Make sure you have the appropriate power cord for the region you are traveling to. For information on purchasing, contact your care provider.

Traveling by plane

⚠ WARNING

- Do not use the device with water in the humidifier tub while in transit (eg. on a plane or vehicle) due to the risk of:
  - water spilling into the device
  - the inhalation of water during turbulence.
- Make sure that the humidifier tub is empty before transporting the device.

Your AirSense device may be taken on board as carry-on luggage. Medical devices do not count toward your carry-on luggage limit.

You can use your AirSense device on a plane as it meets the Federal Aviation Administration (FAA) requirements. Air travel compliance letters can be downloaded and printed from www.resmed.com.

When using the device on a plane:

- Make sure the humidifier tub is empty and inserted into your device. The device will not work without the humidifier tub inserted.
- Make sure the device is switched to airplane mode when required by airline staff.

To turn on Airplane mode:

1. From the Home screen, tap **MORE**.
2. Flick down the menu to locate **Airplane Mode**.
3. Tap **Airplane mode** to switch it on.

Troubleshooting

If you have any problems, have a look at the following troubleshooting topics. If you are not able to fix the problem, contact your care provider or ResMed. Do not try to open the device.

General Issues

Problem/possible cause	Solution
Air is leaking from around my mask	
Mask may be fitted incorrectly.	Make sure your mask is fitted correctly. See your mask user guide for fitting instructions, run the mask fit function or refer to the Mask Fit video in the app.
I am getting a dry or blocked nose	
Humidity level may be set too low.	Increase the **Humidity Level.**
I am getting droplets of water on my nose, in the mask and air tubing	
Humidity level may be set too high.	Decrease the **Humidity Level.**
Tube temperature may be too low.	Increase the **Tube Temperature.**
My mouth is very dry and uncomfortable	
Air may be escaping through your mouth.	You may need a chin strap to keep your mouth closed or a full face mask.
My screen is black	
Power may not be connected.	Connect the AC adaptor and make sure the plug is fully inserted.
My humidifier tub is leaking	
Humidifier tub may not be assembled correctly.	Check for damage and reassemble the humidifier tub correctly.
Humidifier tub may be damaged or cracked.	Contact your care provider for a replacement.
My therapy data has not been sent to my care provider	
Wireless coverage may be poor/The no wireless connection icon  is displayed on the top right of the screen.	Make sure that the device is placed where there is coverage (ie, on your bedside table, not in a drawer or on the floor).
	The wireless signal strength icon  indicates good coverage when all bars are displayed, and poor coverage when fewer bars are displayed.
Device may be in Airplane Mode.	Turn off **Airplane Mode.**
Data transfer is not enabled for your device.	Talk to your care provider about your settings.
SmartStart is enabled, but the device does not automatically start when I breathe into the mask	
Breath is not deep enough to trigger SmartStart	To start therapy, take a deep breath in and out through the mask, before breathing normally.
	Press the Start/Stop button located on top of the device.
There is excessive leak	Adjust the mask and headgear
	Air tubing may not be connected properly. Connect firmly at both ends.

Problem/possible cause	Solution
SmartStop is enabled, but does not automatically stop when I remove the mask.	
Incompatible mask being used	Only use equipment recommended by ResMed.
	Contact ResMed or see ResMed.com for more information.
	If you are using a nasal pillows mask with set pressure less than 7 cm H_2O (7 hPa). SmartStop will not work and should be disabled.
	If you are using a conduit mask, SmartStop will not work and should be disabled.

Device Messages

Problem/possible cause	Solution
System fault, refer to user guide, Error 4	
Device may have been left in a hot environment.	Allow to cool before re-use. Disconnect the AC adaptor and then reconnect it to restart the device.
Air filter may be blocked.	Check the air filter and replace it if there are any blockages. Disconnect the AC adaptor and then reconnect it to restart the device.
There may be water in the air tubing.	Empty the water from the air tubing. Disconnect the AC adaptor and then reconnect it to start the device.
All other error messages, for example, System fault, refer to user guide Error X	
An unrecoverable error has occurred on the device.	Contact your care provider. Do not open the device.

General warnings

⚠ WARNING

- Any change or modification to the product is not expressly approved by ResMed and could void the user's authority to operate the device.
- Supplemental oxygen must not be used while smoking or in the presence of an open flame.
- When using the device with an oxygen supply, check the following:
  - Starting therapy – ensure the device is on and blowing air before the oxygen supply is turned on.
  - Stopping therapy – ensure the oxygen supply is turned off first, then the device.

  This will ensure oxygen does not accumulate within the device and create a risk of fire.
- The device has not been tested or certified for use in the vicinity of X-ray, CT or MRI equipment. Do not bring the device within 13 ft (4 m) of X-ray or CT equipment. Never bring the device into an MR (Magnetic Resonance) environment.
- The device should not be used adjacent to or stacked with other equipment. If adjacent or stacked use is necessary, the device should be observed to verify normal operation in the configuration in which it will be used.
- The use of accessories other than those specified for the device is not recommended. These may increase radio frequency energy or be influenced by the interference and result in improper operation.

For any serious incidents that occur in relation to this device, these should be reported to ResMed and the competent authority in your country.

Technical specifications

65W AC adaptor	
AC input range:	100-240v, 50-60Hz, 2.0A 115V, 400Hz, 1.5A (for aircraft use)
DC output:	24 VDC ± 1 VDC, 2.71A
Typical power consumption:	9.0W
Peak power consumption:	71.9W
Class of Equipment:	Class II

Environmental conditions	
Operating temperature:	+41°F to +95°F (+5°C to +35°C) Note: The airflow for breathing produced by this therapy device can be higher than the room temperature. Under extreme ambient temperature conditions (104°F/40°C) the device remains safe.
Operating humidity:	10 to 95% relative humidity, non-condensing
Operating altitude:	Sea level to 9,870' (3,010 m); air pressure range 700 hPa to 1060 hPa;
Storage pressure/ Storage Altitude	700 to 1060 hPa
Storage and transport temperature:	-13°F to +158°F (-25°C to +70°C)
Storage and transport humidity:	5 to 95% relative humidity, non-condensing

Flow measurement tolerance

± 6 L/min or 10% of reading, whichever is greater, at 0 to 150 L/min positive flow

Pressure measurement tolerance

± 0.5 cm H_2O ±4% of measured reading

Flow (maximum) at set pressures

The following are measured accordingly to ISO 80601-2-70 201.12.1.103

Pressure	AirSense 11, humidifier tub and Standard air tubing	AirSense11, humidifier tub and SlimLine	AirSense 11, humidifier tub and ClimateLineAir 11
cm H_2O (hPa)	L/min	L/min	L/min
4	150	145	144
8	147	142	141
12	143	138	138
16	140	135	134
20	136	131	129

Notes:

- Refer to the relevant measurement uncertainty from the table below
- Values are after gas standard conversion to STPD (ie, correction factor of 0.9962) and additional 5% guard banding to buffer the claimed values and rounded down.

Mode pressure range	
CPAP	4-20 cm H_2O (4-20 hPa) (measured at the mask)
CPAP with EPR mode:	4-20 cm H_2O (4-20 hPa) CPAP with EPR settings: EPR off, Level 1 = 1.0 cm H_2O, Level 2 = 2.0 cm H_2O, Level 3 = 3.0 cm H_2O EPR reduces the pressure during expiration by the amount dependent on the level set above, but the pressure delivered will not drop below 4.0 cm H_2O.
AutoSet, AutoSet for Her mode:	4-20 cm H_2O (4-20 hPa)
AutoSet, AutoSet for Her mode with EPR:	4-20 cm H_2O (4-20 hPa) APAP with EPR settings: EPR off, Level 1 = 1.0 cm H_2O Level 2 = 2.0 cm H_2O,*Level 3 = 3.0 cm H_2O EPR reduces the pressure during expiration by the amount dependent on the level set above, but the pressure delivered will not drop below 4.0 cm H_2O.

Sound

Pressure level measured according to ISO 80601-2-70:2015 and ISO 80601-2-79:2018:

Device with SlimLine and humidifier tub (tub half filled) : 27 dBA with uncertainty of 2 dBA

Power level measured according to ISO 80601-2-70:2015 and ISO 80601-2-79:2018:

Device with SlimLine and humidifier tub (tub half filled): 35 dBA with uncertainty of 2 dBA

Physical - device and humidifier tub

Dimensions (H x W x D):	5.45" x 10.21" x 3.72" (138.5 mm x 259.4 mm x 94.5 mm)
Air outlet:	The 22 mm conical outlet connector complies with EN ISO 5356-1:2015
Wight (device and standard humidifier tub):	40 oz (1130 g)
Weight (device and cleanable humidifier tub):	40 oz (1130 g)
Housing construction:	Flame retardant engineering thermoplastic
Hot plate - material:	Stainless steel
Water capacity:	380 mL
Standard humidifier tub - material:	Injection molded plastic, stainless steel and silicone seal
Cleanable humidifier tub - material:	Injection molded plastic, stainless steel and silicone seal

Air filter

Standard:	Material: Polyester non woven fiber Average arrestance: >75%, when tested to EN779.
Hypoallergenic:	Material: Blended synthetic fibers in a polypropylene carrier Efficiency: >80% (av) when tested to EN13274-7. Note: The use of a ResMed approved hypoallergenic filter will result in a small reduction in the accuracy of the delivered pressure at high leaks.

Design life

Device, power supply unit:	5 years
Cleanable humidifier tub:	2.5 years
Standard humidifier tub	6 months
Air tubing:	6 months

Electromagnetic compatibility

The AirSense 11 complies with all applicable electromagnetic compatibility requirements (EMC) according to IEC 60601-1-2:2014, for residential, commercial and light industry environments.

Portable and mobile RF communications equipment should be used no closer to any part of the machine, including cables, than the recommended 3.94" (10 cm) separation distance.

The AirSense11 has been designed to meet EMC standards. However, should you suspect that the device performance (eg. pressure or flow) is affected by other equipment, move the device away from the possible cause of interference.

Information regarding the electromagnetic emissions and immunity of this ResMed device can be found in ResMed.com/downloads/devices.

IEC 60601-1 (Edition 3.1) classification

Class II (double insulation), Type BF, Ingress protection IP22.

Aircraft use

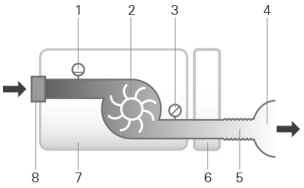
ResMed confirms that the machine meets the Federal Aviation Administration (FAA) requirements (RTCA/DO-160, section 21, category M; RTCA-DO-160, section 20, category T) for all phases of air travel.

Maximum single fault steady state pressure

Device will shut down in the presence of a single fault if the steady state pressure exceeds:

40 cm H_2O (40 hPa) for more than 1 second.

Pneumatic flow path



1. Flow sensor
2. Blower
3. Pressure sensor
4. Mask
5. Air tubing
6. Humidifier
7. Device
8. Inlet filter

Displayed values - Pressure at Mask

	Range	Accuracy	Display resolution
Displayed mask pressure	4-20 cm H_2O	±0.5 cm H_2O ±4% of measured reading	0.1 cm H_2O

Pressure accuracy

Maximum static pressure variation at 10 cm H_2O (10 hPa) according to ISO 80601-2-70:2015

Device with humidifier tub and air tubing:	±0.5 cm H_2O

Note: Refer to the relevant measurement uncertainty from the table below.

Stability of dynamic airway pressure accuracy (short term accuracy) according to ISO 80601-2-70:2015

Device with humidifier tub and air tubing	10 BPM	15 BPM	20 BPM
SlimLine	0.5 cm H_2O	0.7 cm H_2O	1.0 cm H_2O
Standard	0.5 cm H_2O	0.7 cm H_2O	0.8 cm H_2O
ClimateLineAir 11	0.5 cm H_2O	0.7 cm H_2O	1.0 cm H_2O

Measurement system uncertainties

In accordance with ISO 80601-2-70:2015 the measurement uncertainty of the manufacturer's test equipment is:

For measures of flow	± 3.9 L/min
For measures of static pressure	± 0.15 cm H_2O (hPa)
For measures of dynamic pressure	± 0.04 cm H_2O (hPa)

Note: ISO 80601-2-70:2015 stated accuracies and test results provided in this manual for these items already include the relevant measurement uncertainty from the table above.

In accordance with ISO 80601-2-74:2017 the measurement uncertainty of the manufacturer's test equipment is:

For measures of humidification output	± 0.5 mg/L BTPS

Bluetooth

Technology used:	Bluetooth Low Energy (BLE)
Connection types:	GATT
Frequency:	2400 to 2483.5 MHz
Max RF power output:	+4 dBm
Operation range:	10 m (Class 2)

Wireless module

Technology used	Frequencies (MHz)	Max RF power output (dBm)
2G	900/ 1800 *	33.0
3G	850/ 900/ 1700/ 1900/ 2100*	23.5
4G LTE Cat 1	700/ 850/ 1700/ 1900*	23.0

*(bands may not be available in all regions)

FCC ID: 2ACHL-AIR114G

IC: 9103A-AIR114G

The AirSense 11 device complies with FCC Rules and Industry Canada rules. Operation is subject to the following two conditions:

1. This device may not cause harmful interference; and

2. This device must accept any interference received, including interference that may cause undesired operation.

The AirSense 11 device should be installed and operated with minimum distance of 0.59" (15 mm) between the equipment and the user's body.

Additional information regarding the FCC Rules and IC compliance for this device can be found on ResMed.com/downloads/devices

In Canada, the device has been designed to comply with safety standards to radio waves (SAR) in accordance to RSS-102.

Temperature

Maximum heater plate temperature:	154ºF (68ºC)
Temperature Cut-out (heater):	165ºF (74ºC)
Maximum gas temperature: (at mask)[1]	≤ 106ºF (41ºC)
Recommended water type to use in the humidifier tub (Standard tub)	Distilled water (Americas only)
time between each refill of the humidifier tub	> 8 hours ± 0.5 hours*

[1] The air flow for breathing produced by this therapy device can be higher than the temperature of the room. Under extreme ambient temperature conditions (104ºF/40ºC) the device remains safe.

* tested at 23 ±2°C

General

The patient is an intended operator.

Humidifier performance:

SlimLine/Standard tubing

Mask Pressure cm H_2O (hPa)	Nominal % RH when delivered to mask at 72ºF (22ºC)	Nominal % RH delivered to mask, tested in at 72ºF (22ºC) ambient	Nominal system output mg/L AH[1], BTPS[2]	
	Setting 4	Setting 8^ (maximum setting)	Setting 4	Setting 8 (maximum setting)
4	80%	100%	≥6	≥12
10	80%	100%	≥6	≥12
20	80%	100%	≥6	≥12

Climate Control Auto[2] - ClimateLineAir 11

Mask Pressure cm H_2O (hPa)	Nominal % RH delivered to mask, tested in 72ºF (22ºC) ambient	Nominal system output mg/L AH[1], BTPS[2]
4	85%	≥ 12
10	85%	≥ 12
20	85%	≥ 12

[1] AH - Absolute Humidity in mg/L

[2] BTPS - Body Temperature Pressure Saturated

Air tubing

	ClimateLineAir 11	SlimLine/ Standard
ClimateLineAir 11 temperature pressure range	60 to 86ºF (16 to 30ºC)	-
ClimateLineAir 11 temperature cut out	106ºF (41ºC)	106ºF (41ºC)
Maximum recommended pressure	25 cm H_2O	25 cm H_2O
Maximum working temperature, when used with a humidifier	-	106ºF (41ºC)
Material	Flexible plastic and electrical components	Flexible plastic
Inner diameter	0.6" (15 mm)	SlimLine: 0.6" (15 mm) Standard: 0.74" (19 mm)
Length	6'6" (2.0m)	SlimLine: 6' (1.8m) Standard: 6"6' (2.0m)

Note: The manufacturer reserves the right to change these specifications without notice.

Resistance to flow

The table illustrates the resistance to flow of the air tubing:

Air tubing	At flow (L/min) with pressure of 20 cm H_2O	Resistance to flow (cm H_2O/L/min) Air tubing only
Standard	30	0.0050
	15	0.0040
SlimLine	30	0.007
	15	0.006
ClimateLineAir	30	0.0100
	15	0.0077

Compliance

The table illustrates the compliance of the air tubing:

Air tubing	Compliance (cm H_2O/L/min) with pressure of 60 cm H_2O Air tubing only
Standard (19mm) with length of 3m	1.056
SlimLine	0.454
ClimateLineAir	0.500

Guidance and manufacturer's declaration electromagnetic emissions and immunity

⚠ WARNING

Portable RF communications equipment (including peripherals such as antenna cables and external antennas) should be used no closer than 3.9" (10 cm) to any part of the device. Otherwise, degradation of the performance of this equipment could result.

Medical electrical equipment needs special precautions regarding EMC and needs to be installed and put into service according to EMC information provided in this document.

The AirSense device has been designed to meet EMC standards. However, should you suspect that the device performance (eg, pressure or flow) is affected by other equipment, move the device away from the possible cause of interference.

Guidance and manufacturer's declaration - electromagnetic emissions

The device is intended for use in the electromagnetic environment specified below. The patient or the user of the device should assure that the device is used in such an environment.

Emissions test	Compliance	Electromagnetic environment — guidance
RF emissions CISPR 11	Group 1	The device uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emissions CISPR 11	Class B	The device is suitable for use in all establishments, including domestic establishments and those directly connected to the public low-voltage network that supplies buildings used for domestic purposes.
Harmonic Emissions IEC 61000-3-2	Class A	
Voltage Fluctuations/Flicker Emissions IEC 61000-3-3	Complies	

Guidance and manufacturer's declaration - electromagnetic immunity

The device is intended for use in the electromagnetic environment specified below. The customer or the user of the device should assure that the device is used in such an environment.

Immunity test	IEC60601-1-2 test	Compliance level	Electromagnetic environment — guidance
Electrostatic discharge (ESD) IEC 61000-4-2	±8 kV contact ±15 kV air	±8 kV contact ±15 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%.
Electrical fast transient/burst IEC 61000-4-4	±2 kV for power supply lines ±1 kV for input/output lines	±2 kV ±1 kV for input/output lines	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	±1 kV differential mode ±2 kV common mode	±1 kV differential mode ±2 kV common mode	Mains power quality should be that of a typical commercial or hospital environment.
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	<5% Ut (>95% dip in Ut) for 0.5 cycle 70% Ut (30% dip in Ut) for 25 cycles <5% Ut (>95% dip in Ut) for 5 sec	<5% Ut (>95% dip in Ut) for 0.5 cycle 70% Ut (30% dip in Ut) for 25 cycles <5% Ut (>95% dip in Ut) for 5 sec	Mains power quality should be that of a typical commercial or hospital environment. If the user of the device requires continued operation during power mains interruptions, it is recommended that the device be powered from an uninterruptible power source.

Immunity test	IEC60601-1-2 test	Compliance level	Electromagnetic environment — guidance
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	3 A/m	30 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.
Conducted RF IEC 61000-4-6	6 Vrms 150 kHz to 80 MHz	6 Vrms 150 kHz to 80 MHz	Portable and mobile RF communications equipment should be used no closer to any part of the device, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.
Radiated RF IEC 61000-4-3	9 V/m to 28 V/m at frequencies up to 5.785 GHz	27 V/m to 85 V/m at frequencies up to 5.785 GHz	The device complies with all applicable electromagnetic compatibility requirements (EMC) according to IEC60601-1-2:2014, for residential, commercial and light industry environments. Portable and mobile RF communications equipment should be used no closer to any part of the device including cables, than the recommended 3.9" (10 cm) separation distance. The device has been designed to meet EMC standards. However, should you suspect that the device performance (eg pressure or flow) is affected by other equipment, move the device away from the possible cause of interference. Operation is subject to the following two conditions: This device may not cause harmful interference, and this device must accept any interference received including interference that may cause undesired operation.

Notes:

- Ut is the AC mains voltage prior to application of the test level.
- At 80 MHz and 800 MHz, the higher frequency range applies.
- These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

Symbols

Follow instructions before use. Temperature limitation. Humidity limitation. Operating altitude. Atmospheric pressure limitation. Manufacturer. Direct current. On / Off. Class II equipment. IP22 Protected against finger sized objects and against dripping water when tilted up to 15 degrees from specified orientation. Non-ionising radiation. MR unsafe (do not use in the vicinity of an MRI device). RTCA/DO-160 Section 21, Category M Compliant & FAA Compliant. Type BF applied part. Date of Manufacture MD Medical device. REF Catalog number. DN Device number. SN Serial number. LOT Batch code. EC REP European Authorized Representative.

See symbols glossary at ResMed.com/symbols.

Environmental information

This device should be disposed of separately, not as unsorted municipal waste. To dispose of your device, you should use appropriate collection, reuse and recycling systems available in your region. The use of these collection, reuse and recycling systems is designed to reduce pressure on natural resources and prevent hazardous substances from damaging the environment.

If you need information on these disposal systems, please contact your local waste administration. The crossed-bin symbol invites you to use these disposal systems. If you require information on collection and disposal of your ResMed device please contact your ResMed office, local distributor or go to ResMed.com/environment.

California Perchlorate Information:

The coin-cell battery within this device may contain Perchlorate Material - special handling may apply.

See: www.dtsc.ca.gov/hazardouswaste/perchlorate

Servicing

The AirSense device is intended to provide safe and reliable operation when operated in accordance with the instructions provided by ResMed. ResMed recommends that the AirSense device be inspected and serviced by an authorized ResMed Service Centre if there is any sign of wear or concern with device function. Otherwise, service and inspection of the products generally should not be required during their design life.

Limited warranty

ResMed Pty Ltd (hereafter 'ResMed') warrants that your ResMed product shall be free from defects in material and workmanship from the date of purchase for the period specified below.

Product	Warranty period
• Mask systems (including mask frame, cushion, headgear and tubing)—excluding single-use devices • Accessories—excluding single-use devices • Flex-type finger pulse sensors • Humidifier water tubs	90 days
• Batteries for use in ResMed internal and external battery systems	6 months
• Clip-type finger pulse sensors • CPAP and bilevel device data modules • Oximeters and CPAP and bilevel device oximeter adapters • Humidifiers and humidifier cleanable water tubs • Titration control devices	1 year
• CPAP, bilevel and ventilation devices (including external power supply units) • Battery accessories • Portable diagnostic/screening devices	2 years

This warranty is only available to the initial consumer. It is not transferable.

During the warranty period, if the product fails under conditions of normal use, ResMed will repair or replace, at its option, the defective product or any of its components.

This limited warranty does not cover: a) any damage caused as a result of improper use, abuse, modification or alteration of the product; b) repairs carried out by any service organization that has not been expressly authorized by ResMed to perform such repairs; c) any damage or contamination due to cigarette, pipe, cigar or other smoke; and d) any damage caused by exposure to ozone, activated oxygen or other gasses.

Warranty is void on product sold, or resold, outside the region of original purchase.

Warranty claims on defective product must be made by the initial consumer at the point of purchase.

This warranty replaces all other expressed or implied warranties, including any implied warranty of merchantability or fitness for a particular purpose. Some regions or states do not allow limitations on how long an implied warranty lasts, so the above limitation may not apply to you.

ResMed shall not be responsible for any incidental or consequential damages claimed to have resulted from the sale, installation or use of any ResMed product. Some regions or states do not allow the exclusion or limitation of incidental or consequential damages, so the above limitation may not apply to you.

This warranty gives you specific legal rights, and you may also have other rights which vary from region to region. For further information on your warranty rights, contact your local ResMed dealer or ResMed office.

Visit ResMed.com for the latest information on ResMed's Limited Warranty.

Further information

If you require additional information on how to setup, use or maintain the Air11™ system (including ClimateLineAir 11 heated tubing), or to report unexpected operation or events, please contact the ResMed Service Centre or your care provider.

ResMed

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ResMed Pty Ltd
1 Elizabeth Macarthur Drive Bella Vista NSW 2153 Australia

ResMed.com

ResMed

AirSense™ 11

ENGLISH

FCC information (4G)

This document provides additional information regarding compliance of the AirSense 11device with FCC Rules and Industry Canada Rules.

FCC ID: 2ACHL-AIR114G

This device complies with Part 15 of the FCC Rules. Operation is subject to the following two conditions:

- This device may not cause harmful interference,
- This device must accept any interference received, including interference that may cause undesired operation.

This device is designed and manufactured not to exceed the emission limits for exposure to radio frequency (RF) energy set by the Federal Communications Commission of the U.S. Government and Industry Canada.

The exposure standard employs a unit of measurement known as the Specific Absorption Rate, or SAR. The SAR limit relevant for the application described in the manual is 1.6W/kg. Tests for SAR are conducted using standard operating positions accepted by the FCC with the device transmitting at its highest certified power level in all tested frequency bands. Although the SAR is determined at the highest certified power level, the actual SAR level of the equipment while operating can be well below the maximum value. This is because the device is designed to operate at multiple power levels so as to use only the power required to reach the network. In general, the closer you are to a wireless base station antenna, the lower the power output.

Equipment Authorization has been granted to this model with the reported SAR level(s) evaluated as in compliance with the FCC RF exposure guidelines. SAR information on this equipment is on file with the FCC and can be found under the Display Grant section of www.fcc.gov/oet/ea/fccid after searching on the FCC ID as printed on the equipment.

This device has been tested to comply with FCC and IC radiation exposure limits set forth for an uncontrolled environment when used for the documented intended purpose and when mounted and operated as shown in the user guide. The device has been tested for a minimum distance of 15 mm between the device and the human body.

This device complies with the FCC radiation exposure limits set forth in an uncontrolled environment.This equipment has been tested and found to comply with the limits for a Class B digital device, pursuant to part 15 of the FCC Rules. These limits are designed to provide reasonable protection against harmful interference in a residential installation. This equipment generates, uses and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation. If this equipment does cause harmful interference to radio or television reception, which can be determined by turning the equipment off and on, the user is encouraged to try to correct the interference by one or more of the following measures:

- Reorient or relocate the receiving antenna,
- Increase the separation between the equipment and receiver,
- Connect the equipment into an outlet on a circuit different from that to which the receiver is connected,
- Consult the dealer or an experienced radio/TV technician for help.

Any change or modification to the product is not expressly approved by ResMed and could void the user's authority to operate the device.

Industry Canada, IC: 9103A-AIR114G

This device complies with Industry Canada Rules. Operation is subject to the following two conditions:

- This device may not cause interference,
- This device must accept any interference, including interference that may cause undesired operation.

The device has been designed to comply with safety standards for exposure to radio waves (SAR) in accordance to RSS-102.

FRANÇAIS

Renseignements relatifs à la FCC (4G)

Le présent document fournit des renseignements complémentaires relatifs à la conformité du dispositif

AirSense 11 aux règlements de la FCC et aux règles d'Industrie Canada.

ID FCC : 2ACHL-AIR114G

Cet appareil est conforme à la Section 15 des réglementations de la FCC. Son fonctionnement est soumis aux deux conditions suivantes :

- Cet appareil ne doit pas causer de brouillage préjudiciable, et
- Cet appareil doit accepter tout brouillage subi, même si le brouillage est susceptible d'en compromettre le fonctionnement.

Cet appareil est conçu et fabriqué afin de ne pas dépasser les limites d'émission pour une exposition à l'énergie des fréquences radio telles qu'établies par la Commission fédérale des communications du gouvernement des États-Unis (FCC) et par Industrie Canada.

La norme d'exposition est fondée sur une unité de mesure appelée Débit d'absorption spécifique ou SAR (Specific Absorption Rate). Le SAR limite correspondant à l'utilisation décrite dans le manuel est de 1,6 W/kg. Les tests portant sur le SAR sont effectués en utilisant les positions de fonctionnement standard acceptées par la FCC lorsque l'appareil fonctionne à son niveau de puissance certifié le plus élevé dans toutes les bandes de fréquences testées. Bien que le SAR soit déterminé au niveau de puissance certifié le plus élevé, le niveau de SAR réel de l'appareil en cours d'utilisation peut être nettement inférieur à la valeur maximale. Cela est dû au fait que l'appareil est conçu pour fonctionner à différents niveaux de puissance de manière à n'utiliser que la puissance nécessaire pour atteindre le réseau. Généralement, plus vous êtes proche d'une antenne de station de base sans fil, plus la puissance de sortie est faible.

La FCC a accordé une autorisation de commercialisation de ce modèle d'appareil, confirmant la conformité des niveaux SAR évalués avec les directives de la FCC relatives à l'exposition aux RF. Les renseignements relatifs au SAR pour cet appareil sont conservés par la FCC et se trouvent dans la section Display Grant (Afficher l'autorisation) du site www.fcc.gov/oet/ea/fccid après avoir entré l'ID de la FCC apparaissant sur l'appareil dans le champ de recherche.

Cet appareil a été testé et respecte les limites d'expositions aux rayonnements établies par la FCC et IC pour un environnement non contrôlé lorsqu'il est utilisé aux fins prévues, ainsi que monté et utilisé tel que décrit dans le guide de l'utilisateur. L'appareil a été testé pour une distance minimale de 15 mm entre l'appareil et le corps humain.

Cet appareil est conforme aux limites d'exposition au rayonnement de la FCC comme indiquées pour un environnement non contrôlé.

Cet appareil a été testé et respecte les exigences d'un appareil numérique de classe B, conformément à la Section 15 des réglementations de la FCC. Ces limites sont conçues pour procurer une protection raisonnable à l'égard du brouillage préjudiciable dans le cadre d'une installation résidentielle. Cet équipement génère, utilise et peut rayonner de l'énergie radiofréquence et peut causer un brouillage préjudiciable des communications radio, s'il n'est pas installé et utilisé selon les instructions. Toutefois, rien ne garantit l'absence d'interférences dans une installation particulière. Si cet équipement cause un brouillage préjudiciable à la réception des radiocommunications ou des programmes de télévision, ce qui peut être vérifié en allumant et en éteignant l'appareil, l'utilisateur devra essayer de corriger la situation d'une de ces façons :

- Réorienter ou déplacer l'antenne de réception.
- Éloigner l'équipement du récepteur.
- Brancher l'appareil dans une prise reliée à un autre circuit que celui qui alimente le récepteur.
- Communiquer avec le distributeur ou avec un technicien radio/TV expérimenté pour obtenir de l'aide.

Des changements ou des modifications non expressément approuvés par ResMed peuvent annuler le droit d'usage de l'appareil de l'utilisateur.

Industrie Canada, IC : 9103A-AIR114G

Le présent appareil est conforme aux règles d'Industrie Canada. L'exploitation est autorisée aux deux conditions suivantes :

- Cet appareil ne doit pas produire de brouillage, et
- L'utilisateur de l'appareil doit accepter tout brouillage radioélectrique subi, même si le brouillage est susceptible d'en compromettre le fonctionnement.

Cet appareil a été conçu pour respecter les normes de sécurité relatives aux ondes radioélectriques (DAS) conformément à la norme RSS-102.

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