PHILIPS

*Dream*Station Go

Auto CPAP
CPAP

User manual

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1. Safety Information

Intended use

The Philips Respironics DreamStation Go systems deliver positive airway pressure therapy for the treatment of Obstructive Sleep Apnea in spontaneously breathing patients weighing over 30 kg (66 lbs.). It is for use in the home or hospital/institutional environment.

Important

CPAP therapy should only be used on the instruction of a licensed physician.

Several accessories are available to make your Obstructive Sleep Apnea (OSA) treatment with the DreamStation Go system as convenient and comfortable as possible. To ensure that you receive safe and effective therapy, use only Philips Respironics accessories.

⚠ **Warning:** Use only the cleaning methods outlined in your user manual. Philips is unable to verify safety or performance of any device if ozone or other unapproved cleaning and disinfection methods are used.

Note

Any damage caused by unapproved ozone cleaning and disinfection methods or other unapproved cleaning and disinfection methods will not be covered by the Philips Limited Warranty.

⚠ Warnings

A warning indicates the possibility of injury to the user or operator.

Device usage	This device is not intended for life support.
General	Contact your health care professional if symptoms of sleep apnea recur.
Cleaning and Disinfection	Use only the cleaning methods outlined in your user manual. Philips is unable to verify the safety or performance of any device if ozone or other unapproved cleaning and disinfection methods are used. Any damage caused by unapproved ozone cleaning and disinfection methods or other cleaning and disinfection methods will not be covered by the Philips limited warranty.
Improperly functioning device	If you notice any unexplained changes in the performance of the device, if it is making unusual sounds, if water is spilled into the enclosure, or if the enclosure is cracked or broken, discontinue use and contact your health care provider.

Personnel qualifications	This manual serves as a reference. The instructions in this manual are not intended to supersede your health care professional's instructions regarding the use of the device.
	The operator should read and understand this entire manual before using the device.
Operating and storage temperatures	Do not use this device if the room temperature is warmer than 95˚ F (35˚ C) because the temperature of the airflow may exceed 109˚ F (43˚ C). This could cause thermal irritation or injury to the patient's airway.
	Do not use the device while positioned in a warm place, such as direct sunlight or near a heating appliance. These conditions can increase the temperature of the airflow and could cause thermal irritation or injury to the patient's airway.
Bacteria filter	If the device is used by multiple persons, a low-resistance, main flow bacteria filter should be installed in-line between the device and the circuit tubing to prevent contamination.
	If the device is used on multiple users, discard and replace the bacteria filter each time the device is used on a different person.
Patient circuits and tubing	The device should be used only with masks and connectors recommended by Philips Respironics or with those recommended by the health care professional or respiratory therapist. A mask should not be used unless the device is turned on and operating properly. The exhalation port(s) associated with the mask should never be blocked. **Explanation of the Warning:** The device is intended to be used with special masks or connectors that have exhalation ports to allow continuous flow of air out of the mask. When the device is turned on and functioning properly, new air from the device flushes the exhaled air out through the mask exhalation port. However, when the device is not operating, enough fresh air will not be provided through the mask, and exhaled air may be rebreathed.
	If you are using a full face mask (a mask covering both your mouth and your nose), the mask must be equipped with a safety (entrainment) valve.
	Do not pull or stretch the tubing. This could result in circuit leaks.
	Inspect the tubing for damage or wear. Discard and replace the tubing as necessary.
Choking	This device contains small parts which could result in a choking hazard.

Oxygen	When using oxygen with this system, the oxygen supply must comply with local regulations for medical oxygen.
	Do not connect the device to an unregulated or high pressure oxygen source.
	When using oxygen with this system, a Philips Respironics pressure valve must be placed in-line with the patient circuit between the device and the oxygen source. The pressure valve helps to prevent the back flow of oxygen from the patient circuit into the device when the unit is off. Failure to use the pressure valve could result in a fire hazard.
	Oxygen supports combustion. Oxygen should not be used while smoking or in the presence of an open flame.
	Do not use the device in the presence of a flammable anesthetic mixture in combination with oxygen or air, in the presence of nitrous oxide, or in an oxygen-enriched environment.
	Do not use the device near a source of toxic or harmful vapors.
	When using oxygen with this system, turn the device on before turning on the oxygen. Turn the oxygen off before turning the device off. This will prevent oxygen accumulation in the device. **Explanation of the Warning**: When the device is not in operation and the oxygen flow is left on, oxygen delivered into the tubing may accumulate within the device's enclosure. Oxygen accumulated in the device enclosure will create a risk of fire.
Care and Maintenance	Periodically inspect the device, battery pack, electrical cords, cables, tubing, optional humidifier, accessories, and all circuit parts (filter, tube and mask) for damage (such as cracks, tears, or broken pieces) or signs of wear. Discontinue use and replace any damaged parts.
	Repairs and adjustments must be performed by Philips Respironics-authorized service personnel only. Unauthorized service could cause injury, invalidate the warranty, or result in costly device damage. Contact your health care provider for maintenance.
	Do not attempt to modify the device or battery pack in any way.
	Periodically check battery pack charge status and recharge if depleted.
	To avoid electrical shock, always unplug the power cord from the wall outlet before caring for the device. DO NOT immerse the device in any fluids.
	Do not submerge the battery pack in water or any other liquid.

Power cord	Route the power cord to the outlet in a way that will prevent the cord from being tripped over or interfered with by chairs or other furniture.
	This device is activated when the power cord is connected.
	Use only power cords supplied by Philips Respironics for this device. Use of power cords not supplied by Philips Respironics may cause overheating or damage to the device.
	To avoid strangulation hazards, ensure that all cords connected to the device and battery pack are properly routed.
	Periodically inspect the therapy device power cord for signs of wear or damage. If it becomes worn or damaged, contact Philips Respironics or your health care provider for a replacement.
Accessories	Do not use any accessories, detachable parts, and materials not recommended by Philips Respironics. Incompatible parts or accessories can result in degraded performance.
	The USB charging port is designed only for use in charging a mobile device, such as a cell phone. Ensure there are no additional accessories attached to the mobile device while connected to this charging port.
	Use of accessories, transducers and cables other than those specified or provided by the manufacturer of this equipment could result in increased electromagnetic emissions or decreased electromagnetic immunity of this equipment and result in improper operation.
EMC	Use of this equipment adjacent to or stacked with other equipment should be avoided because it could result in improper operation.
	The Health Industry Manufacturers Association recommends that a minimum separation of six inches be maintained between a wireless phone and a pacemaker to avoid potential interference with the pacemaker. The DreamStation Go on-board *Bluetooth* communication should be considered a wireless phone in this regard.
	Do not use this device near active high frequency surgical equipment and the Radio Frequency shielded room of a Medical Electrical system for magnetic resonance imaging, where the intensity of electromagnetic disturbances is high.

Cautions

A caution indicates the possibility of damage to the device.

EMC	Medical electrical equipment needs special precautions regarding EMC and needs to be installed according to EMC information. Contact your health care provider regarding EMC installation information.

Tobacco use	Tobacco smoke may cause tar build-up within the device, which may result in the device malfunctioning.
Mobile RF Communications	Portable RF communications equipment (including peripherals such as antenna cables and external antennas) should be used no closer than 30 cm (12 in.) to any part of the [ME EQUIPMENT or ME SYSTEM], including cables specified by the manufacturer. Otherwise, degradation of the performance of this equipment could result.
Device usage	Before operating the device, ensure that both end caps are attached whenever any of the accessories such as the battery pack is not installed.
	Ensure that the therapy device is properly secured if it is being used in a portable environment.
Electrostatic Discharge (ESD)	Pins of connectors shall not be touched and connections shall not be made without special precautions. Precautionary procedures include methods to prevent build-up of electrostatic charge (e.g., air conditioning, humidification, conductive floor coverings, non-synthetic clothing), discharging one's body to the frame of the equipment or system or to earth. It is recommended that all individuals that will handle this device understand these precautionary procedures at a minimum as part of their training.
Condensation	Condensation may damage the device. If the device has been exposed to either very hot or very cold temperatures, allow it to adjust to room temperature (operating temperature) before starting therapy. Do not operate the device outside of the operating temperature range shown in the Specifications section later in this manual.
Care and Maintenance	Allow the device, battery pack, and humidifier to dry completely before reconnecting to the power source.
Filters	A properly installed, undamaged Philips Respironics reusable filter or disposable, fine filter is required for proper operation.
	Clogged inlet filters may cause high operating temperatures that may affect device performance. Regularly examine the inlet filters as needed for integrity and to check for accumulated debris.
	Never install a wet filter into the device. You must ensure sufficient drying time for the rinsed filter.
	Make sure the air inlet holes on the side of the device are not blocked by bedding, curtains, or other items. Air must flow freely around the device for the system to work properly.

Battery Pack	Do not expose the battery pack to extreme temperatures (see the Specifications section for temperature specifications). If the battery pack becomes very hot or very cold, allow it to come to room temperature before using.
	There are no user-serviceable parts inside of the battery pack; therefore, do not attempt to disassemble or repair it.
Extension cords	Do not use extension cords with this device.
Device placement	Do not place the device in or on any container that can collect or hold water.
	Do not place the device directly onto carpet, fabric, or other flammable materials.

Notes

- Any serious incident that has occurred in relation to this device should be reported to Philips and the competent authority of the Member State in which the user and/or patient is established.
- An electronic copy of these instructions can be found at: www.philips.com/IFU.

Contraindications

When assessing the relative risks and benefits of using this equipment, the clinician should understand that this device can deliver pressures up to 20 cm H_2O. In the event of certain fault conditions, a maximum pressure of 40 cm H_2O is possible. Studies have shown that the following pre-existing conditions may contraindicate the use of CPAP therapy for some patients:

- Bullous Lung Disease
- Pathologically Low Blood Pressure
- Bypassed Upper Airway
- Pneumothorax
- Pneumocephalus has been reported in a patient using nasal Continuous Positive Airway Pressure. Caution should be used when prescribing CPAP for susceptible patients such as those with: cerebral spinal fluid (CSF) leaks, abnormalities of the cribriform plate, prior history of head trauma, and/or pneumocephalus. (Chest 1989; 96:1425-1426)

The use of positive airway pressure therapy may be temporarily contraindicated if you exhibit signs of a sinus or middle ear infection. Not for use with patients whose upper airways are bypassed. Contact your health care professional if you have any questions concerning your therapy.

Safety Symbols Glossary

The following symbols may appear on the device and accessories:

Symbol	Title and Meaning
	Operator's manual; operating instructions Consult instructions for use.
	Electronic instructions for use Indicates that relevant information for use of the product is available in electronic form.
	For airline use. Complies with RTCA/DO-160G section 21, category M.
	AC power (Alternating current) Indicates on the rating plate that the equipment is suitable for alternating current only; to identify relevant terminals.
	Separate collection for electrical and electronic equipment per EC Directive 2012/19/EU.
	Li Ion Battery General symbol for recovery/recyclable Lithium-Ion battery.
	*Bluetooth*® symbol Indicates the device has *Bluetooth* capabilities.
IP22	Drip proof equipment Protection against ingress of solid foreign objects ≥ 12.5 mm diameter. Protection against ingress of water with harmful effects dripping (15° tilted).
	Non-ionizing electromagnetic radiation Indicates that the equipment includes RF transmitters.
	Caution, consult accompanying documents.
	Electrostatic sensitive devices (ESD warning symbol) Attention – Observe precautions for handling electrostatic sensitive devices.
	Serial connection

Symbol	Title and Meaning
	Class II equipment (Double Insulated) To identify equipment meeting the safety requirements specified for Class II equipment.
	Keep away from sunlight Indicates the medical device needs protection from light sources.
	Type BF applied part To identify a type BF applied part complying with IEC 60601-1.
	Do not disassemble.
	For indoor use only Equipment is designed primarily for indoor use.
	Medical Device Indicates that the item is a medical device.
	MR unsafe Do not use device in a Magnetic Resonance (MR) environment.
	Unique Device Identifier Indicates the Unique Device Identifier information.
	Packaging unit Indicates the number of pieces in the package.
	Single patient use Indicates that the tubing is for single patient use only.
	Date of Manufacture To indicate the date on which a product was manufactured. Country of Manufacturer To indicate the country of manufacture of the product. Note: When applied to the label, “CC” is replaced by the two letter country code.
	Importer Indicates the entity importing the medical device.

2. System Overview

The DreamStation Go CPAP is a Continuous Positive Airway Pressure therapy device designed for the treatment of Obstructive Sleep Apnea (OSA).

Several accessories are also available for use with your device. Contact your health care provider to purchase any accessories not included with your system.

System Contents

Your DreamStation Go system may include the following items:

• Device	• microSD Card (optional)
• User Manual	• Disposable Fine Filter (optional)
• Reusable Filter	• Battery Pack (optional)
• 12 mm Micro-flexible Tubing (12 Type)	• Heated Humidifier (optional)
• 6 ft. (1.83 m) Power Cord	

Note

If any of these items are missing, contact your health care provider.

Note

For information about the heated humidifier, refer to the DreamStation Go Heated Humidifier user manual.

Accessories

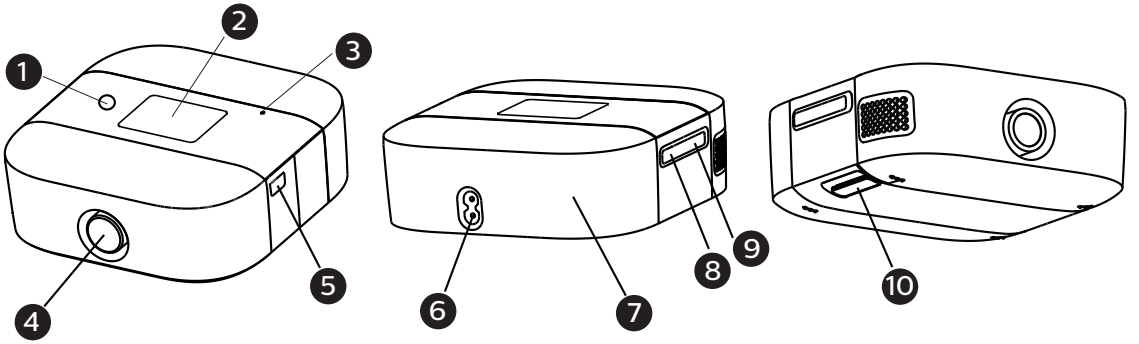
The following accessories are available for your DreamStation Go system:

• 6 ft. (1.83 m) Serial Communication Cable (with ferrites)	• 10 ft. (3.04 m) Power Cord
• Small Travel Kit	• Medium Travel Kit
• 15 mm Standard Tubing (15 Type)	• 22 mm Performance Tubing (22 Type)
• Bacteria Filter	• Pressure Valve (for use with supplemental oxygen)

Note

Your mobile device charging cable should not be longer than 6 ft (1.83 m).

System Diagram



The figure above illustrates some of the device features, described in the following table.

#	Feature	Description
1	Therapy on/off button ◯	Starts and stops the airflow for therapy.
2	Display Touchscreen	This is the User Interface for the therapy device.
3	Ambient Light Sensor	Detects room light levels and adjusts brightness of the display screen.
4	Air Outlet Port	Connect the tubing here.
5	Serial Connector	Access the serial connector here.
6	AC Power Inlet	Connect the power cord here.
7	Battery Pack Access	This end cap slides off for access to the battery pack connection.
8	microSD Card	Access the microSD card here.
9	Mobile Charging Port	Access the USB charging port here for mobile device usage.
10	Filter Access	Access the filter here.

3. Therapy Device

Where to Place Your Device

Place the device on a firm, flat surface somewhere within easy reach of where you will use it. The device should sit at a level lower than your sleeping position. Make sure the device is away from any heating or cooling equipment (e.g., forced air vents, radiators, air conditioners).

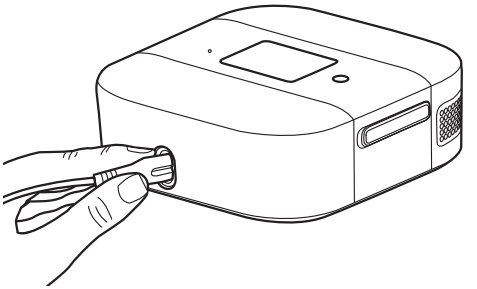
Note

When positioning the device, make sure that the power cable is accessible because removing power is the only way to turn off the device.

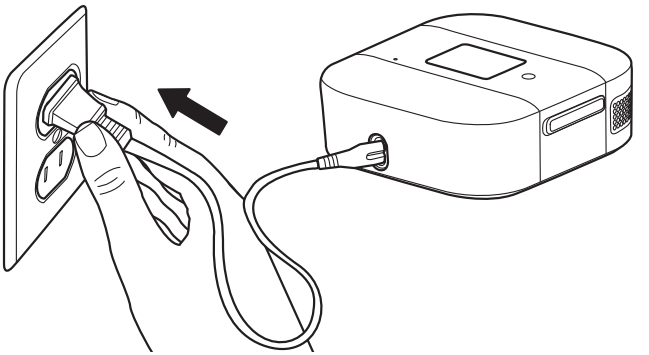
Supplying AC Power to Your Device

Complete the following steps to operate the device by plugging the AC power cord into an electrical outlet.

1. Plug the power cord connector into the power inlet on the back of the device.



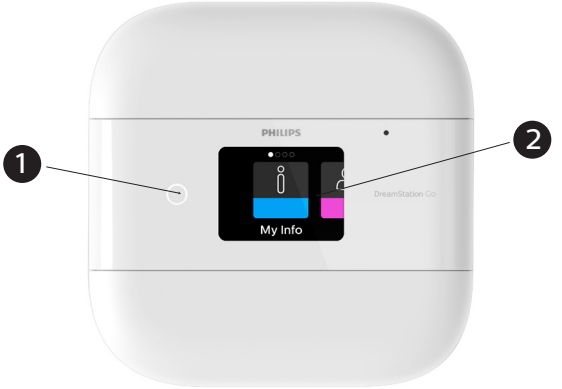
2. Plug the pronged end of the AC power cord into an electrical outlet that is not controlled by a wall switch.



3. Make sure that all connections are secure.

Starting the Device

1. Ensure power is supplied to the device. **Tap anywhere on the display screen to wake up the device.**



#	Feature
1	Therapy on/off button
2	Display Touchscreen

2. Put on your mask assembly. Refer to the instructions supplied with the mask.
3. Tap the therapy button ◯ on top of the device to turn on airflow and begin therapy. The current delivered pressure will display on the screen. The therapy button is **only for therapy.**
4. A small amount of mask leak is normal and acceptable. Correct large amount of mask leaks or eye irritation by adjusting your mask headgear. See the instructions provided with your mask for more information, or refer to the Check Mask Fit section.
5. Tap the therapy button again to turn off the therapy. To turn off therapy when the display screen is off, place and hold a finger on the display screen for three seconds. Alternatively, tap anywhere on the display screen to wake up the display and then tap the therapy button to turn off therapy.

Notes

- If you are using the device in a bed with a headboard, try placing the tubing over the headboard. This may reduce tension on the mask.
- During therapy, if there is a power loss, the device will return to the home screen once power is restored. You may resume therapy as needed.

Navigating the Device Screens

The User Interface (UI) on this device allows you to adjust the device settings and view information about your therapy. The UI consists of the display screen and the touch panel. Swipe left or right on the touch panel to scroll through the menu options on the display screen.

To adjust a setting:

1. Swipe the touchscreen until you find your desired menu option.
2. Tap the desired menu option.
3. Swipe the touchscreen until you find the sub-menu option and tap to select that setting.
4. Swipe the touchscreen to change the setting.
5. Tap the icon or tap the up arrow in the upper left corner of the display to save the setting, and return to the previous menu option.

Note

- The swipe icon on any screen indicates to swipe the display left or right to perform an action.
- The tap icon on any screen indicates to press the display to perform an action.
- Tapping the down arrow on the display when the down arrow appears on any screen will take you to a sub-menu with more menu options. Tapping the up arrow when the up arrow appears on any sub-menu will return you back to the main menu.
- The screens shown throughout this manual are examples for reference only. Actual screens may vary based upon device model and health care provider settings.

Therapy On Menu Navigation Settings

While the device is delivering therapy, you can adjust or view the following settings.

#	Feature	Description
1	Therapy pressure	Displays the current delivered pressure.
2	Ramp feature	If Ramp is enabled, pressing the Ramp icon will return the device to the Ramp set pressure.
3	MyStart Restart	If MyStart is enabled, pressing the Restart icon will return the device to the MyStart set pressure.

Note: The MyStart and Ramp features are not available at the same time. Your health care provider will enable only one.

Ramp Feature

Ramp is a comfort feature that reduces the air pressure you receive when you are trying to fall asleep and then gradually increases (ramps) the pressure until your prescription setting is reached, allowing you to fall asleep more comfortably.

If ramp is enabled on your device, after you turn on the airflow, tap the Ramp ( ) icon on the display. You can use the ramp feature as often as you wish.

When you tap the Ramp icon, the therapy screen will change to reflect the ramp in pressure, and the numbers within the blue circle will reflect the gradual increase in pressure.

Your device has two ramp modes. The standard ramp mode increases pressure at a steady rate. Alternately, the SmartRamp mode maintains a constant lower pressure until the device detects that you require more pressure.

MyStart Feature

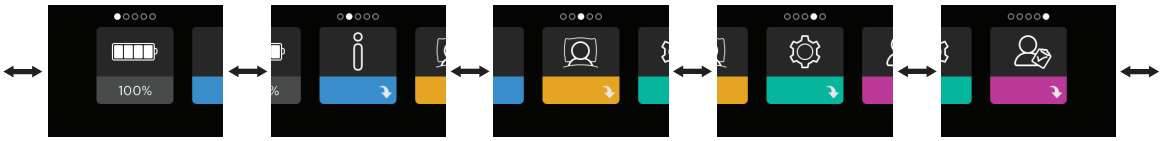
The MyStart feature allows you to choose a starting Auto CPAP pressure at which you are the most comfortable to fall asleep. If your pressure feels too low or too high while trying to fall asleep you can adjust your MyStart pressure up or down from the MyComfort menu.

If MyStart is enabled, the MyStart Pressure is the pressure you will receive when you turn on the airflow until the MyStart Time expires. You can return to your MyStart Pressure at any time during the night by tapping the Restart icon .

When you turn on the airflow or tap the Restart icon, the blue circle on the therapy screen will change to reflect that the time in this feature is approaching the MyStart Time. The numbers within the blue circle will initially reflect the MyStart Pressure. As the night progresses this number will update to reflect any necessary therapy change in pressure between the MyStart Pressure and the device maximum pressure.

Therapy Off Menu Navigation Settings

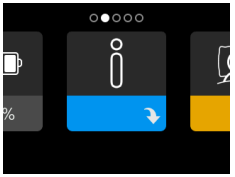
From the Home screen, you can scroll between the following five options:



Battery My Info My Comfort My Device My Support

Battery	This menu is visible when the battery pack is connected. See **Chapter 7, Battery Pack** for details.
My Info	This menu provides summary statistics of your therapy use.
My Comfort	This menu contains comfort settings that you can adjust as needed.
My Device	This menu contains device settings that you can change.
My Support	This menu contains information that your health care provider may direct you to read to them so they can better assist you over the phone.

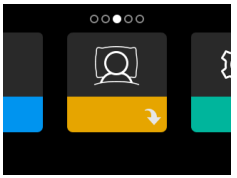
My Info



When you select **My Info**, you will be able to view the following screens. You cannot change settings in the My Info menu. These screens are only for reference. Your health care provider may periodically ask you for this information. If any of the below options are not visible, your health care provider did not enable those options.

Icon	Text	Description
AHI	Last Session	This screen displays the amount of time the user is actually receiving therapy on the device, Apnea/Hypopnea indices (AHI) value and value “100% minus Large Leak” for the most recent 1-day time frame.
	Therapy Hours	This screen displays the amount of time the user is actually receiving therapy on the device for the most recent 1-day time frame. It also displays the average amount of time the patient is actually receiving therapy over the last 7 and 30 days.
AHI	AHI	This screen displays the Apnea/Hypopnea indices (AHI) value for the most recent 1 day time frame. It also displays the average of these individual AHI values over a 7 and 30 day time frame.
	Mask Fit	Displays the value “100% minus Large Leak”. Large Leak is the percentage of time that the mask leak was so high that it is no longer possible for the device to identify respiratory events with statistical accuracy. Displays the value for the most recent 1 day, as well as the values over last 7 and 30 days.
CSR	Periodic Breathing	Displays the percentage of time that the user experienced periodic breathing. Displays the value for the most recent 1 day time frame, as well as values for the last 7 and 30 days. If you observe a large increase in the percent of time in periodic breathing indicated here, contact your health care provider for assistance.
90%	90% Pressure	This screen displays the value of 90% Pressure for the most recent 1 day time frame, as well as values for the last 7 and 30 days.

My Comfort

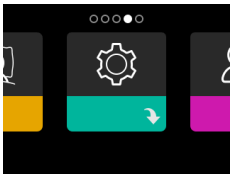


When you select **My Comfort**, you will be able to view the following screens. You can change the settings in the setup menu. These screens will only display if they are available and enabled on your device. If any of the below options are not visible, your health care provider did not enable those options. If a lock icon is displayed on this screen, it indicates that your provider has locked this setting and you cannot change it.

Icon	Text	Description
	Ramp	This displays the ramp starting pressure. You can increase or decrease the ramp starting pressure in 0.5 cm H_2O increments.
	Ramp Time	When you set the Ramp Time, the device increases the pressure from the value set on the ramp screen to the therapy pressure setting over the length of time specified here.
	MyStart Pressure	This displays the current MyStart Pressure. You can adjust the pressure from 4.0 to 20.0 cm H_2O in 0.5 cm H_2O increments.
	MyStart Time	When you set the MyStart Time, the device will override your minimum therapy pressure over the length of time specified here.
	Flex (Type)	This allows you to select the type of air pressure relief that you feel when you exhale during therapy from the available Flex types on your device. You may also turn off this feature.
	Flex (Level)	This allows you to adjust the level of air pressure relief that you feel when you exhale during therapy. Your health care provider can enable or disable this feature. When your health care provider enables Flex, a level will already be set for you on the device. You can increase or decrease the setting from 1 to 3. The setting of “1” provides a small amount of pressure relief, with higher numbers providing additional relief.
	Mask Type	If using a Philips Respironics mask with your DreamStation Go, check the mask’s instructions for use for a “System One” resistance control value (e.g., X1, X2). If one exists, set the “System One” resistance control value that is specified for your Philips Respironics mask for optimal pressure relief (Flex) delivery. Contact your health care provider if you cannot find the resistance setting for your mask.

Icon	Text	Description
	Tube Type	This setting allows you to select the correct tubing type that you are using with the device. You can choose (12) for the Philips Respironics 12 tubing type, (15) for the Philips Respironics 15 tubing type, or (22) for the Philips Respironics 22 tubing type. Note: The 12 type and 15 type tubing are identified on the cuff with the tubing identifier symbol: “12” or “15”. The 22 tubing type does not have any identifier on the cuff.
	Check Mask Fit	This feature allows you to check the fit of your mask prior to starting therapy. This is done by measuring the amount of leak.

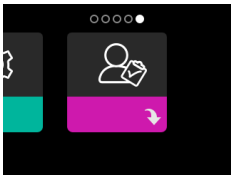
My Device



When you select **My Device**, you will be able to view the following screens. You can change the settings in the setup menu. These screens will only display if they are available and enabled on your device. If any of the below options are not visible, your health care provider did not enable those options.

Icon	Text	Description
	Therapy Ring	This setting controls the therapy button LED light ring during therapy. The LED light ring will remain on during therapy if you select Light On. The LED light ring will fade with the display backlight if you select Light Dims.
	Language	This feature allows you to choose which language to display on the interface.
	*Bluetooth*	This feature allows you to turn *Bluetooth* off and on. Also, it allows you to clear the pairing with a compatible *Bluetooth* device.
	Time	This setting allows you to adjust the time. The default setting is Greenwich Mean Time, but you may adjust the time in 30 minute increments to match your local time zone. **Note: This time setting should not be used as a clock function on the device. It is to align your therapy data for your health care provider’s data reports.**

My Support



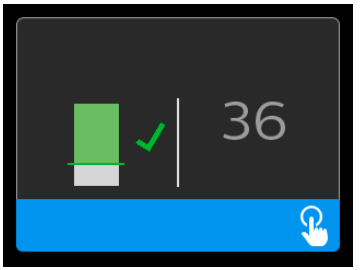
When you select **My Support**, you will be able to view the following screens. You cannot change settings in the support menu. These screens are only for reference. Your health care provider may periodically ask you for this information. If any of the below options are not visible, your health care provider did not enable those options.

Icon	Text	Description
	Device (Info)	This screen displays your therapy device information: serial number, model, and software version.
	Therapy (Info)	This screen displays therapy information your health care provider may request to support you.
	Phone-In	This screen displays the total therapy hours and total blower hours for the device, and a compliance check number used by your health care provider to validate that data provided by you is the data taken from this screen.
	Performance (Check)	Your device is equipped with a self-diagnostic tool called “Performance Check.” This tool can evaluate your device for certain errors. It also allows you to share key device settings with your health care provider. Use Performance Check when directed to by your health care provider. At conclusion of the scan, the screen displays a green check mark if no issue is detected. If device displays a red “X,” please contact your health care provider for assistance.
A-Trial	A-Trial	If Auto-Trial mode is available, this screen displays Days: xx/xx (where xx/xx is the number of completed trial days/ number of selected trial days).

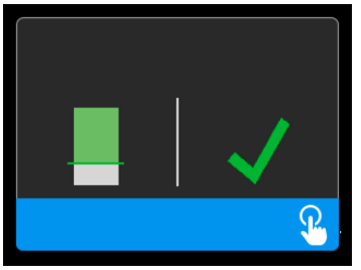
Check Mask Fit

The optional **Check Mask Fit** feature can be enabled or disabled by your health care provider. This feature allows you to check the fit of your mask prior to starting therapy.

1. Put on your mask assembly. Refer to your mask instructions if needed.
2. Navigate to the **Check Mask Fit** screen under **My Comfort** and tap the display to initiate the check.
3. The device will deliver a test pressure while the screen counts down 40 seconds.



4. After the test is complete, the screen will either display a green check mark or a red "X". The green checkmark indicates that there is an appropriate amount of leak. The red "X" indicates that the leak may affect device performance, however, the device will remain functional and deliver therapy.



Note

If you choose to try to improve your mask fit, you can stop therapy, adjust the fit of your mask, and rerun the **Check Mask Fit** feature. Refer to the instructions that came with your mask and headgear for the proper fitting procedure.

Pairing Therapy Device to *Bluetooth*®-enabled Mobile Device

Your device may have *Bluetooth* wireless technology, which is one method by which you can transfer your therapy device's data to **DreamMapper**. DreamMapper is a mobile and web-based system designed to help you enhance your sleep therapy experience.

Notes

- You can only pair your therapy device to one mobile device at any given time.
- Pairing works best when your therapy device and mobile device are in the same room.
- The current version of DreamMapper will guide you through these instructions.
- After initiating pairing, you will have 30 seconds to complete the setup. After this time, it will be cancelled automatically.

Follow the steps below to manually pair to your mobile phone or tablet.

1. Install DreamMapper on your mobile device.
2. With your therapy device powered up and the blower off, initiate *Bluetooth* Setup from the DreamMapper mobile app.
3. The therapy device will appear as **PR BT XXXX** (XXXX will be the last four digits of the serial number listed on the bottom of your therapy device or in **My Support** settings).
4. Your mobile device will require you to confirm pairing via one of these two methods:

— *Enter a PIN code*

The following icon will appear on your therapy device screen with **Pair?**: 

Swipe left or right to select "yes," and tap the display to confirm your setting. Your therapy device will display a 6 digit PIN. Enter this PIN on your mobile device to complete pairing.

— *Confirm a PIN code*

The following icon will appear on your therapy device screen with a 6-digit PIN and **Pair?**:

Verify that the PIN is the same on both the therapy device and the mobile device. If so, swipe the therapy device's display to select "yes" and tap the display to select. Then, accept on the mobile device to complete pairing.

Device Pop-Up Messages

Device pop-ups are messages that show up on the user interface screen. Additional pop-up messages are contained in each chapter.

The following summary table summarizes the messages:

Condition	Icon	Description	Possible Cause	Action
Time		Prompts to set the time.	n/a	Set the time on the device.
MyStart Pressure Adjust		Set the MyStart Pressure setting.	n/a	Adjust your MyStart Pressure and tap the display to save and clear the message.
Sleep Progress	n/a	Shows a three night summary of therapy.	n/a	Tap the display to acknowledge and clear the message.
Change Accepted		Confirms acceptance of prescription change or device upgrade.	n/a	Tap the display to acknowledge and clear the message.
EZ-Start Pressure Incremented to xx.x		Displays when EZ-Start is enabled and device is increasing therapy pressure therapy setting for the next session.	n/a	No action needed.
Pair?: 123456 Yes/No		Prompts to accept or decline pairing to a *Bluetooth* compatible device. This device can be identified by the digits displayed.	n/a	Swipe the display to accept pairing (Yes), or decline (No), then tap the display to confirm selection. The pop-up will timeout after 30 seconds and the pairing will be cancelled if you do not select Yes.

3. Therapy Device

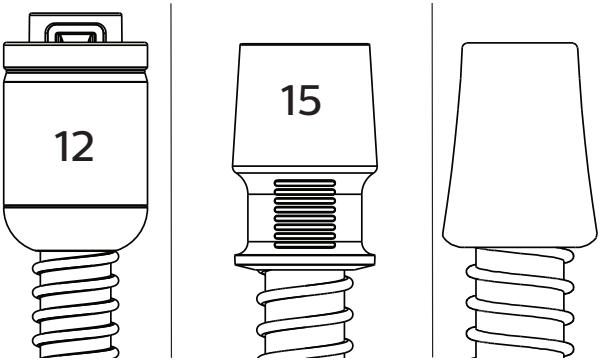
Condition	Icon	Description	Possible Cause	Action
Bluetooth LE Passkey		Prompts to accept or decline pairing to a *Bluetooth* compatible device before displaying the pairing passkey.	n/a	If you selected Yes to accept pairing, the *Bluetooth* LE Passkey will display a passkey on the screen. Enter the passkey on your mobile device to pair. The pop-up will timeout after 30 seconds and the pairing will be cancelled if you do not use the passkey.
Patient Message		Message from your health care provider.	n/a	Tap the display to acknowledge and clear the message.
Change Rejected		A prescription or settings change was rejected.	Change missing or incorrect.	Contact your health care provider.
Service Required		Indicates an error which enters device into “Safe State.” This allows power to remain on but airflow is disabled.	Device error	Disconnect device from power. Reattach power cord to restore power. If the alert continues to occur, contact your provider.
Automatic Off		Displayed when therapy ends due to automatic off function.	The mask has been removed.	Put your mask back on, confirm good fit, and turn airflow on to resume therapy.
Loading Language and Rebooting		Displayed when a new language is selected from the menu.	n/a	No action needed. Times out when complete.

Condition	Icon	Description	Possible Cause	Action
Busy		Displayed when the device is temporarily inaccessible due to data communication.	n/a	No action needed.
Software Upgrade		Prompts to update the device for software changes.	n/a	Choose between “Yes”/”No” when asked to upgrade the software. If “yes” is selected, the upgrade will be made. Do not remove from power. If you select “no”, the message will be cleared.

4. Tubing

Tube Types

There are three different types of tubing that you may use with your DreamStation Go therapy device. You must select the tube type on your device.



12 tubing type | **15 tubing type** | **22 tubing type**

The 12 tubing type will have a “12” identified on the tubing cuff (as shown in the image above). The 15 tubing type will have a “15” identified on the tubing cuff (as shown in the image above). The 22 tubing type does not have any number or symbol on the tubing cuff (as shown in the image above).

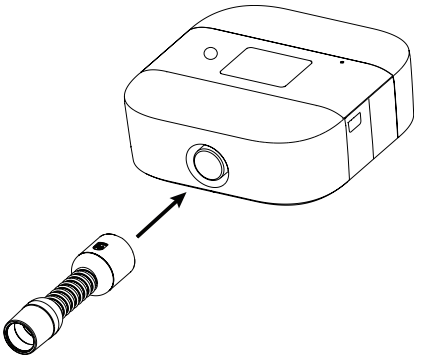
Connecting the Tubing to Your Device

To connect the tubing to your device, you will need the following Philips Respironics accessories:

- Nasal or full face mask (interface) with built-in exhalation, or a nasal or full face mask (interface) with a separate exhalation device attached (such as the Whisper Swivel II)
- Flexible tubing, 6 ft. (1.83 m)
- Mask headgear

Follow these steps to connect tubing to your device:

1. Insert the 12, 15 or 22 tubing type cuff into the air outlet port on your therapy device.



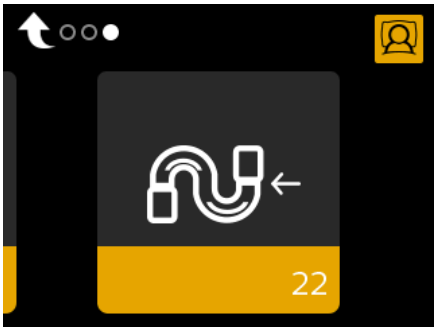
2. Connect the tubing to your mask. For proper placement and positioning, refer to the instructions that came with your mask.

Notes

- Use the supplied mask adapter if your mask does not directly connect to the 12 tubing type.
- You may use a standard tube with a bacteria filter. If required, connect a bacteria filter to the device air outlet, and then connect the flexible tubing to the outlet of the bacteria filter. When using the bacteria filter, the device performance may be affected. However, the device will remain functional and deliver therapy.
- The mask connection port (cuff) of the 12, 15 and 22 mm tubing types is 22 mm per standard ISO 5356-1.

Changing Your Tube Type

Change your tube type by navigating to **My Comfort -> Tube Type**. Swipe left or right to switch between tube types.



4. Tubing

For more information on navigation or selection, please refer to **Chapter 3, Navigating the Device Screens** or **My Comfort**.

Tubing Device Pop-Up Messages

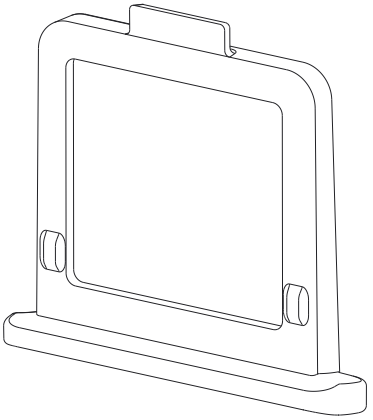
Device pop-ups are messages that show up on the user interface screen.

Condition	Icon	Description	Possible Cause	Action
Low leak: Check mask and tube		Blocked airway	Blockage at tube or mask	Check tube is not crushed or folded, such that airflow is restricted. Check mask is attached properly and without any obstruction.

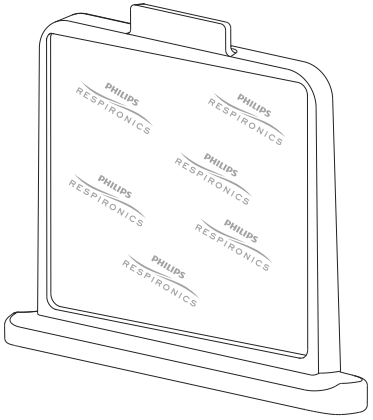
5. Filter

Filter Types

You may use either a reusable filter that is rinseable, or a disposable, fine filter.



Reusable Filter



Disposable Fine Filter

The **reusable filter** screens out normal household dust and pollen. The reusable filter is supplied with your device. Rinse the reusable filter weekly and dispose of it monthly.

The **disposable, fine filter** provides more complete filtration of fine particles. The disposable, fine filter is recommended for people who are sensitive to tobacco smoke or other small particles. The disposable, fine filter is sold separately. The disposable, fine filter contains Philips Respironics branding in the media (shown in the image above).

DO NOT rinse the disposable, fine filter. Dispose of it monthly.

When using the disposable, fine filter, the device performance may be affected. However, the device will remain functional and deliver therapy.

Installing or Replacing the Filter

One of the filters **must be in place at all times** to operate the device. If a filter is not already installed in the device, you must at least install the reusable filter before using your device.

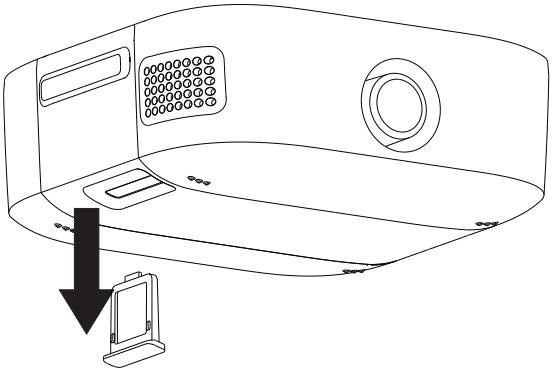
The device has an automatic air filter reminder. Every 30 days, your device will display a message reminding you to check your filters and replace them as needed.

Note

The air filter reminder is a message only. The device does not detect the performance of the filters, nor does it recognize when a filter has been replaced.

Follow these steps to install/replace a filter into your device:

1. If replacing an existing filter, pull out the old filter assembly.



2. Insert a dry, reusable filter or a new, disposable fine filter into the filter access on the device.

Filter Device Pop-Up Messages

Device pop-ups are messages that show up on the user interface screen.

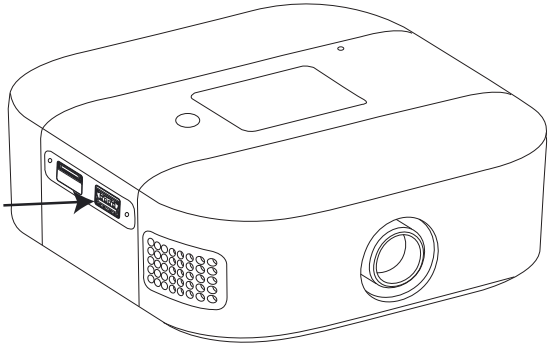
Condition	Icon	Description	Possible Cause	Action
Inlet blocked. Check filter.		Blocked airway	Blockage at device inlet.	Check device air inlet is not obstructed. Check air filter(s) are installed properly; replace if needed.

6. Accessories

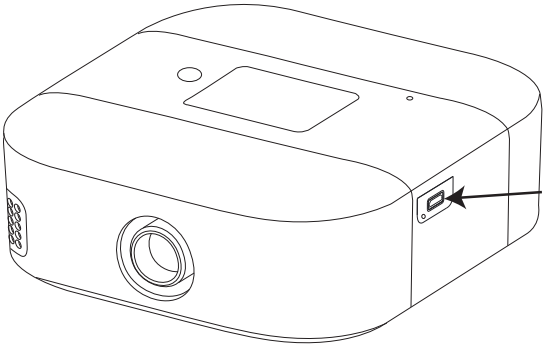
There are several accessories available for your DreamStation Go system, such as a microSD card, a travel kit or a battery pack. The device also comes with a USB port and a micro USB port. The travel kit is available for convenient portability while traveling with your device. Contact your health care provider for additional information on the available accessories. When using optional accessories, always follow any instructions enclosed with the accessories.

Using the USB Port and the Micro USB Port

The DreamStation Go device comes with a USB port and a micro USB port. The USB port may be used to charge your mobile devices. The micro USB port may be used by your health care provider to extract therapy data. Remove the cover over each port to access.



USB Port



Micro USB Port

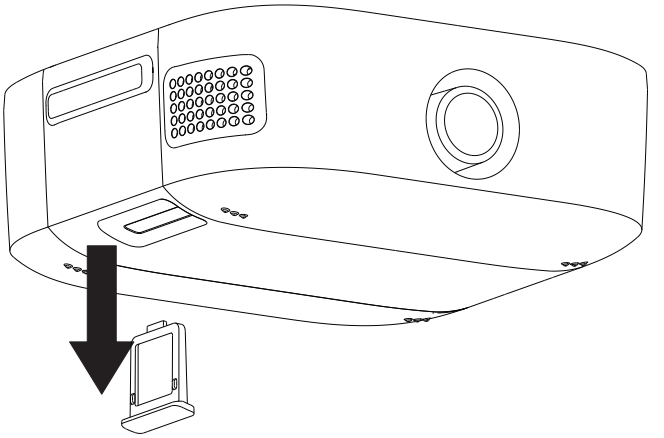
Using the microSD Card

The DreamStation Go system may come with a microSD card inserted in the microSD card slot on the side of the device to record information for your health care provider. Your health care provider may ask you to periodically remove the microSD card and send it to them for evaluation.

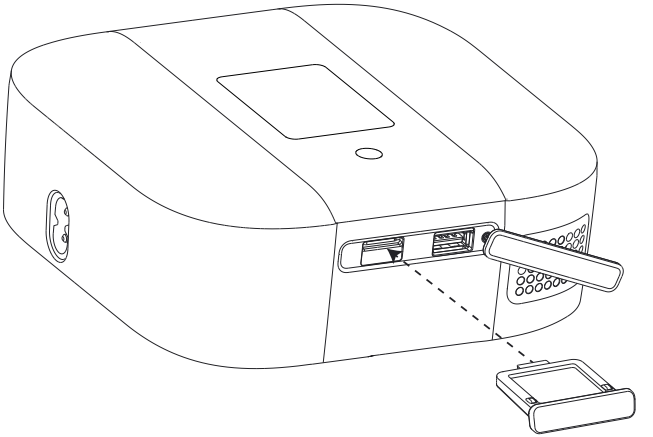
You will use the filter to remove the microSD card.

Turn off therapy and follow these steps to remove the microSD card:

1. Remove the filter from the device. Refer to the **Installing or Replacing the Filter** section in Chapter 5 of this manual.



2. Use the end of the filter to push in on the microSD card. This will push the microSD card out of the device.



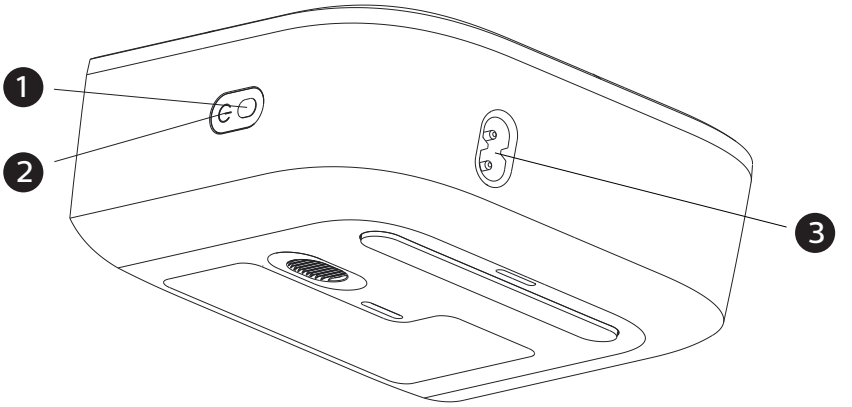
microSD Card Device Pop-Up Messages

Device pop-ups are messages that show up on the user interface screen.

Condition	Icon	Description	Possible Cause	Action
Data Activity: Do not remove microSD card		microSD card read/ write underway.	n/a	No action needed. Message will clear when the microSD card activity is finished.
microSD card removed		Indicates microSD card has been removed from therapy device and not reinserted before the start of the current therapy session.	microSD card was not reinserted into device.	Reinsert microSD card, or click to clear alert.
microSD card error: Remove and reinsert		microSD card error detected.	Device cannot read the microSD card. A problem may exist with the microSD card or it was ejected during a writing activity, or it was inserted incorrectly.	Remove microSD card and reinsert. If message reappears, contact your health care provider for a replacement card.
microSD card full		microSD card is full.	microSD card is full.	Remove microSD card and replace with a new card from your health care provider.

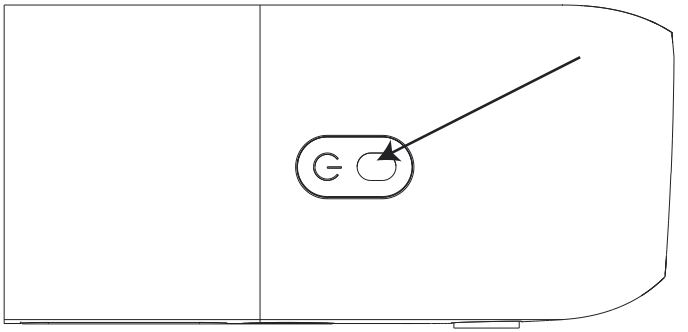
7. Battery Pack

Indicators and Buttons on the Battery Pack



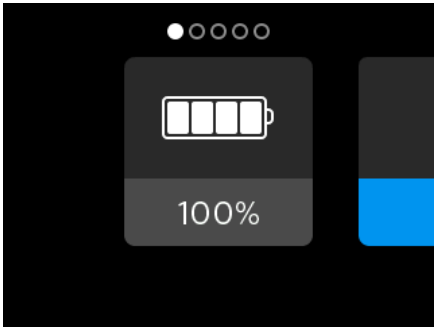
#	Feature
1	Push Button
2	LED Display
3	AC Power Inlet

- Push Button - The push button is located on the LED display of the battery pack.
- LED Display - The battery pack uses one green LED light to indicate the battery pack charge status when the battery pack is charging while not connected to the therapy device (standalone charging). The LED will be in one of the three modes:
  * Steady ● when the battery pack is fully charged
  * Blinking ✳ when charging
  * Off ○ when connected to the therapy device



- Battery Pack Charge Indicator During Therapy – The display touchscreen shows the current battery pack charge status in the lower right hand corner when the pack is connected and therapy is active (shown below). A fully charged battery pack is indicated by four charge bars. The white charge bars disappear as the battery pack charge decreases. Depending on your settings, mask leak and environmental conditions, a fully charged battery pack typically lasts 8 hours. For further information, speak with your health care provider.

- Battery Pack Charge Indicator When Therapy Is Not Active – The display touchscreen will show the battery charge percentage when the battery is connected to the therapy device but not in use (shown below). This screen will appear in your main menu selections.



- Battery Pack State of Charge Alert – The display touchscreen will display a battery with a question mark in the center (shown below) when the charge level cannot be determined.

7. Battery Pack

- Battery Pack Fault Alert - The display touchscreen will display a battery with an X inside (shown below) when a battery fault is detected.

Preparing the Battery Pack for First Use and Recharging

1. Remove the battery pack from the packaging.
2. Plug the end of the AC power cord into the battery pack.
3. Plug the AC power cord into an AC outlet. The battery pack will begin to charge automatically.
4. Once the battery pack is fully charged, it is ready for use with the DreamStation Go therapy device.

Notes

- Periodically charge the battery pack if not used regularly.
- Retain your packaging in case you ever need to return your battery pack to Philips Respironics.
- Before using the battery pack for the first time, you must plug it in until it is fully charged. This may take up to 5 hours.

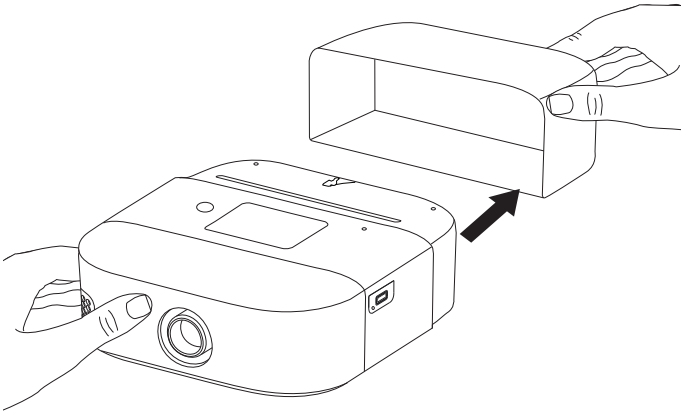
Attaching the Battery Pack to the Device

After charging, your battery pack is ready to use. It can either be disconnected from the AC outlet and used as an external battery pack (standalone mode), or remain plugged into the AC outlet for a continuous, fully charged battery pack (uninterruptible power supply (UPS) mode).

To use the battery pack in UPS mode, follow the below steps:

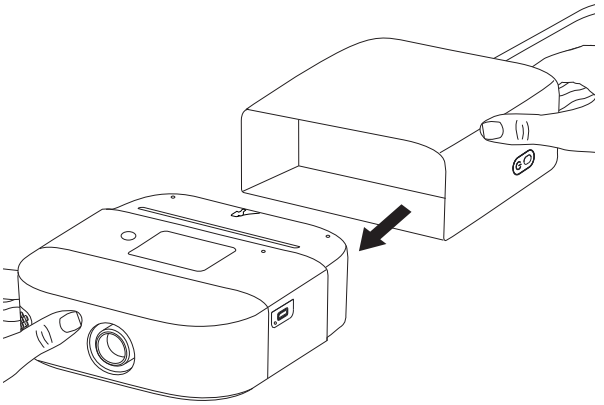
1. Keep the battery pack plugged into the power source and connected to an AC outlet. This will allow you to use the battery pack continuously without losing any charge.

2. Remove the battery pack end cap on the therapy device.



3. Slide the battery pack onto the device where the end cap was. Make sure the battery pack latches onto the therapy device.

4. Attach the AC power cord to the battery pack and then to the AC outlet.



To use the battery pack in standalone mode, follow these steps:

1. Make sure the battery pack is fully charged. Disconnect the power cord from the AC outlet and disconnect the power cord from the therapy device. It can now be used with your therapy device as an external battery pack.

2. Remove the battery pack end cap on the therapy device.

3. Slide the battery pack onto the device where the end cap was. Make sure the battery pack latches onto the therapy device.

4. Momentarily push the battery pack push button to wake up the battery pack.

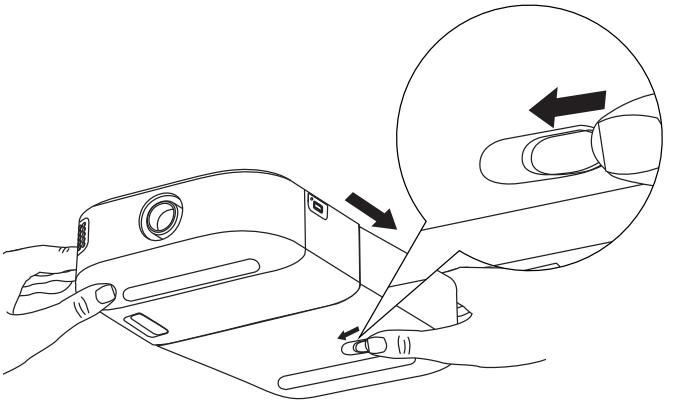
7. Battery Pack

Notes

- The first time you charge your battery pack, it must be **fully charged** in accordance with the **Preparing the Battery Pack for First Use and Recharging** section. After the first charge is complete, the battery pack will charge while connected in UPS mode.
- To preserve battery life in UPS mode, the battery pack stops charging when it reaches full charge. The battery will start charging again when it depletes to 90% charge status.
- When the battery pack is used in standalone mode and the therapy device enters standby mode, the therapy device automatically shuts down the battery pack to preserve battery charge.

Disconnecting the Battery Pack

1. Disconnect the power cord.
2. Hold the battery pack push button down for 5 seconds, or the battery pack will shut off within 30 minutes when not in use.
3. The power down pop-up message will appear and the therapy device will power down and go dark.
4. You can now disconnect the battery pack. Disengage the battery pack by sliding the latch on the back of the pack, and pulling the battery pack away from the therapy device.



5. Replace the battery pack end cap on the therapy device.

8. Cleaning, Disinfection & Maintenance

Cleaning the CPAP Device and Optional Battery Pack

Inspect your device and optional battery pack weekly to see if they need care.

1. Turn the device off and disconnect from the power source. Remove any cables attached to the device or battery pack.
2. Remove the reusable filter. Refer to "Caring for the Reusable Filter" below for more information.
3. Wipe for 1 minute with a lint-free cloth dampened with a cleaning solution (1 teaspoon [5 ml] of mild liquid dish washing detergent [such as Dawn® Ultra Dishwashing Liquid][1] per gallon [3.8 liters] of potable water) to clean the exteriors of the therapy device and the battery pack (if used).
4. Pay close attention to all corners and crevices of the device's exterior surfaces. Be sure all visible soil is removed.
5. Wipe with a lint-free cloth dampened (not dripping) with potable water for 1 minute, turning the cloth frequently to remove all detergent residue.
6. Let everything air dry completely.
7. After cleaning, inspect the device, battery pack, and all circuit parts (filter, tube, and mask) for damage such as cracks, tears, or broken pieces. If any parts are damaged, contact Philips Respironics Customer Service. Replace any damaged parts.

Caution

Allow the device and battery pack to dry completely before reconnecting to the power source.

Note

The expected service life of the base device is 5 years.

Disinfecting the Exterior of the CPAP Device & Optional Battery Pack

In the hospital environment, clean and disinfect the exterior of the device weekly and between each patient.

Before disinfecting the CPAP device and battery pack, be sure that it has been cleaned as instructed in the previous section of this guide.

1. After cleaning, use a lint-free cloth dampened with 70% isopropyl alcohol to wipe the alcohol onto the exterior, thoroughly wetting all surfaces.
2. Keep wet for 4 minutes.
3. Allow to air dry.
4. Inspect the device for damage after cleaning and disinfection. If any parts are damaged, contact customer service.

1 Dawn is a registered trademark of Procter & Gamble.

Caution

Allow the device and battery pack to dry completely before reconnecting to the power source.

Note

The expected service life of the base device is 5 years.

Cleaning the Tubing and Adapters

In the home environment, for single-patient use, clean tubing and any adapters before first use and weekly. Replace tubing and adapters annually.

1. Disconnect the flexible tubing from the device.
2. Gently wash the tubing, including any adapters or connectors, for 3 minutes by completely immersing in a cleaning solution (1 teaspoon [5 ml] of mild liquid dish-washing detergent [such as Dawn Ultra Dishwashing Liquid] per gallon [3.8 liters] of potable water).

   Note: During immersion, gently move the tubing back and forth to loosen and adequately remove adhering substances from the tubing, adapters, and connectors.

   Note: Be sure to clean the entire inner surface of the tube by ensuring it is fully immersed in the detergent solution during gentle agitation by hand.
3. Rinse thoroughly with potable water for at least 1 minute to remove all soap residue from the tubing, adapters, and connectors and allow to air dry completely. Make sure all parts are dry before next use.
4. Visually inspect the tubing for cleanliness. Repeat the cleaning if not visually clean.
5. Inspect the tubing for damage or wear (cracking, tears, punctures, etc.). Discard and replace if necessary.

Note

Replace tubing after 6 months.

Caring for the Reusable Filter

In the home environment, for single-patient use, rinse the reusable filter **at least once a month** and replace it with a new one every **six months**.

The disposable, fine filter should be replaced after 30 nights of use, or sooner if it appears clogged. **DO NOT rinse the fine filter.**

Follow these steps to rinse the reusable filter:

1. If the device is operating, stop the airflow. Disconnect the device from the power source.
2. Remove the filter from the device. Refer to the **Installing or Replacing the Filter** section in **Chapter 5**.
3. Take the reusable filter to a sink, turn it upside down (tabs down), and run tap water through the white middle portion of the filter to rinse away any debris.
4. Shake the filter to remove as much water as possible.
5. **Allow the filter to air dry completely before reinstalling it.** If the filter is damaged, replace it.
6. Reinstall the filter into the filter access area on the device. Refer to the **Installing or Replacing the Filter** section in **Chapter 5**.

Notes

- Only Philips Respironics supplied filters should be used as replacement filters.
- If soil is visible during inspection, rinse more frequently.
- Replace the disposable, fine filter if it is damaged or has accumulated debris.

9. Troubleshooting

Tips and Tricks

Your device is equipped with a self-diagnostic tool called **Performance Check**. This tool can evaluate your device for certain errors. It also allows you to share diagnostic information with your health care provider. Use Performance Check when directed by your health care provider.

The table below lists some of the problems you may experience with your device and possible solutions to those problems.

Contact customer service for assistance if none of the below troubleshooting tips work for you.

Problem	Why it happened	What to do
Nothing happens when you apply power to the device. The backlights on the buttons do not light.	There's no power at the outlet or the device is unplugged.	• If you are using AC power: Check the outlet and verify that the device is properly plugged in, and that there is power available at the outlet. Make sure the AC power cord is connected correctly to the device's power inlet. • If you are using the battery pack: Make sure your battery pack is securely connected to your device. If the battery pack has been exposed to extreme temperatures, allow the battery pack to cool or warm to room temperature. Check to see if your battery pack needs charged or replaced.
The airflow does not turn on.	There may be a problem with the blower.	• Make sure the device is powered correctly. • Make sure the home screen appears on the user interface. • Press the therapy button on top of the device to start airflow. If the airflow does not turn on, there may be a problem with your device.

Problem	Why it happened	What to do
The device's display is erratic.	The device has been dropped or mishandled, or the device is in an area with several electronic devices.	Unplug the device. Reapply power to the device. If the problem continues, relocate the device to an area away from electronic equipment (such as cellular phones, cordless phones, computers, TVs, electronic games, hair dryers, etc.).
The Ramp feature does not work when you press the Ramp button.	Your health care provider did not enable Ramp for you, or your therapy pressure is already set to the minimum setting	• If ramp has not been enabled for you, discuss this feature with your health care provider. • If your health care provider has enabled ramp, but the feature still does not work, check the current pressure setting on the therapy screen. If the therapy pressure is set to the minimum setting (4.0 cm H_2O), or the ramp starting pressure is the same as the therapy pressure, the ramp feature will not work. Make sure that the ramp time setting is >0.
The airflow is much warmer than usual.	The air filters may be dirty. The device may be operating in direct sunlight or near a heater.	• Rinse or replace the air filter. • The temperature of the air may vary somewhat based on your room temperature. • Make sure that the device is properly ventilated. Keep the device away from bedding or curtains that could block the flow of air around the device. • Make sure the device is away from direct sunlight and heating equipment.
The airflow pressure feels too high or too low.	The tubing type setting may be incorrect.	Make sure the tubing type setting (12, 15 or 22) matches the tubing that you are using (Philips Respironics 12, 15 or 22 tubing type).

Problem	Why it happened	What to do
I hear a leak or whistling sound coming from my therapy device (not related to mask leak).	The therapy device air inlet may be obstructed.	• Check therapy device air inlet is not obstructed, and filter has not accumulated excessive debris and is properly inserted. • Confirm that the device and tube are connected properly and not leaking.
The battery pack LEDs will not light up while charging.	Your battery pack may have been damaged.	If the battery pack is completely depleted of charge, wait a few minutes for the LEDs to light up. If the LEDs still do not light up, replace your battery pack. If the battery pack has been exposed to extreme temperatures, allow the battery pack to cool or warm to room temperature.
The battery pack LED is rapidly flashing.	Your battery pack may have been damaged.	If the battery pack has been exposed to extreme temperatures, allow the battery pack to cool or warm to room temperature. Unplug the battery pack from the power cord, then plug the power cord back into the battery pack. If the LED continues to rapidly flash, replace your battery pack.
“Service Required” shown on display.	A device error has occurred and placed the device into safe state.	Disconnect power cord. Reattach the power cord to restore power. If the alert continues, contact your health care provider.

Contacting Customer Service

Should you experience trouble with this equipment or require assistance setting up, using, or maintaining the device, please contact your health care provider. If you need to contact Philips Respironics directly, contact customer service at +1-724-387-4000, or go to www.respironics.com to find your local customer service contact information.

10. Additional Notes

Traveling with the System

When traveling, the optional case is for carry-on luggage only. The optional case will not protect the system if it is put through checked baggage.

For your convenience at security stations, there is a symbol on the bottom of the device indicating that it is medical equipment and is suitable for airline use. It may be helpful to bring this manual along with you to help security personnel understand the DreamStation Go device.

If you are traveling to a country with a line voltage different than the one you are currently using, a different power cord or an international plug adaptor may be required to make your power cord compatible with the power outlets of the country to which you are traveling. Contact your health care provider for additional information.

Airline Travel

The device is suitable for use on airlines when the device is operating from an AC power source or battery pack.

Altitude Compensation

This device automatically compensates for altitude up to 7,500 feet. No manual adjustment is necessary.

Adding Supplemental Oxygen

Oxygen can be added to the patient circuit.

Notes

- Refer to the pressure valve's instructions for complete setup information.
- When using oxygen with this system, turn the device on before turning on the oxygen. Turn the oxygen off before turning the device off. This will prevent oxygen accumulation in the device.
- Do not connect the device to an unregulated or high pressure oxygen source.

Service

The device does not require routine servicing.

Additional Notices

Notices:

- The *Bluetooth*® word mark and logos are registered trademarks owned by Bluetooth SIG, Inc. and any use of such marks by Philips Respironics is under license. Other trademarks and trade names are those of their respective owners.
- Dawn® is a registered trademark of Procter & Gamble.
- The DreamStation Go Therapy Device is capable of transmitting data between the therapy device and a mobile device. This connection between the therapy device and a mobile device is encrypted.
- This device contains a FCC certified *Bluetooth* radio module (located on the main board).
- Use of non-original manufacturer-approved accessories may violate your local RF exposure guidelines and should be avoided.
- This device complies with part 15 of the FCC 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. 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, TV reception, or other devices which can be determined by turning the equipment on and off, the user is encouraged to try to correct the interference by one or more of the following measures:

  Reorient or relocate the receiving antenna (on the radio, TV, or other device).

  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 of the device for help.
- A small portion of the firmware that performs data encryption on the DreamStation Go device is being utilized under the Apache 2.0 and Mozilla 2.0 licenses. These licenses are available at: www.apache.org/licenses/LICENSE-2.0 and https://www.mozilla.org/en-US/MPL/2.0/
- Any changes or modifications made to the device that are not expressly approved by Respironics may void the user's authority to operate the equipment.

Hereby, Respironics Inc. declares that this class 1 radio equipment is in compliance with Directive 2014/53/EU. The full text of the EU declaration of conformity is available at the following internet address: http://incenter.medical.philips.com/PMSPublic.

Specifications

Environmental

Operating Temperature	Device: 5° to 35° C (41° to 95° F)
	Battery Pack: 5° to 35° C (41° to 95° F)
Storage Temperature	-20° to 60° C (-4° to 140° F)
Relative Humidity	15 to 95% (non-condensing); (operating & storage)
Atmospheric Pressure:	Device: 101 to 77 kPa (0 - 2286 m / 0 - 7500 ft)
	Battery Pack: 101 to 77 kPa (0 - 2286 m / 0 - 7500 ft)

Physical

Dimensions	150.8 x 150.8 x 58.8 mm (5.937” L x 5.937” W x 2.315” H)	(Device)
	122 x 150.8 x 58.8 mm (4.803” L x 5.937” W x 2.315” H)	(Battery Pack)
Weight (Approximate)	854 g (1.88 lb.)	(Device)
	696 g (1.53 lb.)	(Battery Pack)

Expected Service Life

The expected service life of the DreamStation Go therapy device is 5 years.

The expected service life of the battery pack is 3 years.

Standards Compliance

This device is designed to conform to the following standards:

- IEC 60601-1 General Requirements for Basic Safety and Essential Performance of Medical Electrical Equipment
- IEC 60601-1-11 General Requirements for Basic Safety and Essential Performance in the Home Healthcare Environment
- IEC 60601-1-6 General Requirements for Safety - Usability
- IEC 62366-1 Application of Usability Engineering in Medical Devices
- IEC 62304 Medical Device Software – Software Life-cycle Processes
- ISO 80601-2-70 Sleep Apnea Breathing Therapy Equipment
- EN 60601-1-2 Electromagnetic Compatibility
- RTCA/DO-160G section 21, category M; Emission of Radio Frequency Energy

IEC 60601-1 Classification

Type of Protection Against Electric Shock	Class II Equipment/Internally Powered
Degree of Protection Against Electric Shock	Type BF Applied Part
Degree of Protection Against Ingress of Water	Device: Drip Proof, IP22 First characteristic numeral (2): Protection against ingress of solid foreign objects ≥ 12.5 mm diameter. Explanation: Protected against access to hazardous parts with a finger and protected against solid foreign objects of 12.5 mm diameter and greater. Second characteristic numeral (2): Protection against ingress of water with harmful effects dripping (15° tilted). Explanation: Protected against vertically falling water drops when enclosure tilted up to 15°.
Mode of Operation	Continuous

Electrical

AC Power Consumption	100–240 VAC, 50/60 Hz, 2.0-1.0 A
Fuses	There are no user-replaceable fuses.
USB Charging Port Output	5 VDC, 7.5 W (1.5 A)

Electrical for the Battery Pack

Run Time	> 8 hours
Output voltage range	18-24.6 VDC
Battery technology	Lithium Ion
Capacity	62 Wh
Input voltage range	100-240 VAC, 50/60 Hz, 2.0-1.0 A
Output power (max continuous)	50W
Minimum life cycle	≥ 70% of rated capacity after 500 cycles
Recharge time	< 5 hours

Radio Specifications

Operating Frequency Range	2402 - 2480 MHz
Maximum Output Power	< 10 dBm
Modulation	GFSK, P/4 DQPSK, 8DQPSK

Intake Port Filters

Reusable Filter	100% Polyester 88% Efficient @ 7-10 micron size
Fine Filter	Blended Synthetic Fiber 95% Efficient @ 0.5-0.7 micron size

Declared Dual-number Noise Emissions (in Accordance with ISO 4871)

Tube Size	Sound Pressure Level (L)	Uncertainty (K)	Sound Power Level (L)	Uncertainty (K)
12 (mm) tubing type	32 dB(A)	2 dB(A)	40 dB(A)	2 dB(A)
15 (mm) tubing type	31 dB(A)	2 dB(A)	39 dB(A)	2 dB(A)
22 (mm) tubing type	31 dB(A)	2 dB(A)	39 dB(A)	2 dB(A)

Note

Values determined according to noise test code given in ISO 80601-2-70:2015, using the basic standards ISO 3744 and ISO 4871.

Pressure Accuracy

Pressure Increments: 4.0-20.0 cm H_2O (in 0.5 cm H_2O increments)

Maximum static pressure accuracy, according to ISO 80601-2-70:2015:

Tube Type	Pressure	Accuracy
12 (mm) tubing type	10 hPa (cm H_2O)	± 1.0 hPa (cm H_2O)
15 (mm) tubing type and 22 (mm) tubing type	10 hPa (cm H_2O)	± 0.5 hPa (cm H_2O)

Static pressure accuracy has a measurement uncertainty of 3.8%

Maximum dynamic pressure variation, according to ISO 80601-2-70:2015:

Tube Type	10 BPM	15 BPM	20 BPM
12 (mm) tubing type	0.6 hPa (cm H_2O)	0.8 hPa (cm H_2O)	1.6 hPa (cm H_2O)
15 (mm) tubing type and 22 (mm) tubing type	0.7 hPa (cm H_2O)	0.7 hPa (cm H_2O)	1.0 hPa (cm H_2O)

Dynamic pressure accuracy has a measurement uncertainty of 3.6%.

Maximum Flow Rate (Typical)

Tube Type	Flow	Test pressures (hPa/cm H_2O)				
		4.0	8.0	12.0	16.0	20.0
12 (mm) tubing type	Average flow at the patient connection port (l/min)	90	119	112	106	99
15 (mm) tubing type	Average flow at the patient connection port (l/min)	77	115	112	105	106
22 (mm) tubing type	Average flow at the patient connection port (l/min)	80	121	127	121	109

Tubing Specifications

Tubing	Tubing resistance (RI/RE)	Tubing compliance (@ 60 hPa)
12 mm performance tubing (PR12)	@30 l/min: 0.03 hPa/l/min (cmH2O/l/min)	0.44 ml/hPa (ml/cmH2O)
15 mm performance tubing (PR15)	@15 l/min: 0.006 hPa/l/min (cmH2O/l/min) @30 l/min: 0.01 hPa/l/min (cmH2O/l/min)	0.52 ml/hPa (ml/cmH2O)
22 mm performance tubing	@15 l/min: 0.00 hPa/l/min (cmH2O/l/min) @30 l/min: 0.003 hPa/l/ min (cmH2O/l/min)	0.73 ml/hPa (ml/cmH2O)

Disposal

Dispose of this device in accordance with local collections and recycling regulations. For more information, visit www.philips.com/recycling.

This device contains a lithium ion battery that cannot be recycled. Please contact customer service at +1-724-387-4000, go to www.respironics.com to find your local customer service contact information, or send an e-mail to service@respironics.com for safe disposal instructions of this part.

EMC Information

Your unit has been designed to meet EMC standards throughout its Service Life without additional maintenance. There is always an opportunity to relocate your DreamStation Go Therapy Device within an environment that contains other devices with their own unknown EMC behavior. If you believe your unit is affected by locating it closer to another device, simply separate the devices to remove the condition.

Pressure and Flow Accuracy

The DreamStation Go Therapy Device is designed to perform within the pressure and flowrate accuracies specified in the user manual. If you suspect that the pressure and/or flow rate accuracy is affected by EMC interference remove power and relocate the device to another area. If performance continues to be affected discontinue use and contact your healthcare provider.

Guidance and Manufacturer's Declaration - Electromagnetic Emissions – This device is intended for use in the electromagnetic environment specified below. The user of this device should make sure it 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 power supply network.**
Harmonic emissions IEC 61000-3-2 220~240VAC 50/60Hz (No standard test requirement for this device at other voltage/frequency ranges)	Class A 220~240VAC 50/60Hz	
Voltage fluctuations/Flicker emissions IEC 61000-3-3 220~250VAC 50/60Hz (No standard test requirement for this device at other voltage/frequency ranges)	Complies 220~250VAC 50/60Hz	
Emission of Radio Frequency Energy RTCA/DO-160G Section 21	Category M	This device is suitable for use onboard commercial airplanes inside passenger cabin.

10. Additional Notes

Guidance and Manufacturer's Declaration - Electromagnetic Immunity – This device is intended for use in the electromagnetic environment specified below. The user of this device should make sure it is used in such an environment.

Immunity Test	IEC 60601 Test Level	Compliance Level	Electromagnetic Environment – Guidance
Electrostatic Discharge (ESD) IEC 61000-4-2	±8 kV contact discharges ±2 kV, ±4 kV, ±8 kV, & ±15 kV air discharges	±8 kV contact discharges ±2 kV, ±4 kV, ±8 kV, & ±15 kV air discharges	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 35%.
Electrical fast Transient/burst IEC 61000-4-4	±2 kV for power supply lines and ±1 kV for input-output lines; both at 100 kHz repetition rate.	±2 kV at 100 kHz repetition rate for Power Supply Lines NA - Device does not have user I/O lines that are longer than 3m in length.	Mains power quality should be that of a typical home or hospital environment.
Surge IEC 61000-4-5	±1 kV differential mode ±2 kV common mode	±1 kV differential mode N/A: Device is Class 2 and does not have earth connection.	Mains power quality should be that of a typical home or hospital environment.

Immunity Test	IEC 60601 Test Level	Compliance Level	Electromagnetic Environment - Guidance
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	<5% U_T (>95% dip in U_T) for 0.5 cycle at 45 degree increments <5% U_T (>95% dip in U_T) for 1 cycle 70% U_T (30% dip in U_T) for 0.5 seconds <5% U_T (>95% dip in U_T) for 5 seconds	<5% U_T (>95% dip in U_T) for 0.5 cycle at 45 degree increments <5% U_T (>95% dip in U_T) for 1 cycle 70% U_T (30% dip in U_T) for 0.5 seconds <5% U_T (>95% dip in U_T) for 5 seconds	Mains power quality should be that of a typical home or hospital environment. Upon power interruption, the device may reboot. 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 supply or a battery.
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	30 A/m	30 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical hospital or home environment.
NOTE: U_T is the AC mains voltage prior to application of the test level.			

Guidance and Manufacturer's Declaration - Electromagnetic Immunity – This device is intended for use in the electromagnetic environment specified below. The user of this device should make sure it is used in such an environment.

10. Additional Notes

Immunity Test	IEC 60601 Test Level	Compliance Level	Electromagnetic Environment – Guidance
Conducted RF IEC 61000-4-6 Radiated RF IEC 61000-4-3	3 Vrms 150 kHz to 80 MHz 6 Vrms Amateur Radio & ISM Bands between 150 kHz and 80 MHz 10 V/m 80 MHz to 2.7 GHz Telecommunication frequencies as specified in clause 8.10 of IEC 60601-1-2:2014: 450, 810, 870, 930, 1720, 1845, 1970, and 2450 MHz at 28 V/m 385 MHz at 27 V/m 710, 745, 780, 5240, 5500, and 5785 MHz at 9 V/m	3 Vrms 150 kHz to 80 MHz 6 Vrms Amateur Radio & ISM Bands between 150 kHz and 80 MHz 10 V/m 28 V/m 27 V/m 9 V/m	Portable and mobile RF communications equipment should be used no closer to any part of the device, including cables, than the recommended 30 cm (12 in) separation distance. Interference may occur in the vicinity of equipment marked with the following symbol:

If you plan on using your DreamStation Go on an airplane, please detach this letter and bring it with you.

PHILIPS

FAA Compliance

To Whom It May Concern:

The following Philips Respironics DreamStation Go components are in compliance with commercial airline EMI/RFI requirements:

- DreamStation Go CPAP
- DreamStation Go Auto CPAP
- DreamStation Go Battery Pack

Philips Respironics has designed and tested the identified devices for compliance with section 21, Category M, RTCA DO-160 EMI/RFI requirements as specified in the Code of Federal Regulations 14 CFR 382 *"Nondiscrimination on the Basis of Disability in Air Travel; Final Rule"*.

In accordance with these requirements, the aforementioned components may be used onboard an aircraft without further testing by the carrier.

If you have any other questions regarding our products, please feel free to call the Philips Respironics Customer Services department at 1-724-387-4000. You can also use the following addresses:

Philips Respironics
1001 Murry Ridge Lane
Murrysville, PA 15668

PHILIPS
RESPIRONICS

www.philips.com/respironics

Limited Warranty

Respironics, Inc., a Philips company (“Philips Respironics”) provides this non-transferable, limited warranty for Dreamstation Go (“Product”) to the customer who originally purchased the Product directly from Philips Respironics.

What this warranty covers: Philips Respironics warrants each new Product will be free from defects in materials and workmanship and will perform in accordance with the Product specifications under normal and proper use and maintenance in accordance with applicable instructions, subject to the exclusions below.

This warranty lasts two (2) years for the therapy device and the humidifier (not including the water tank) from the longer of the date of shipment to the purchaser or date of setup by purchaser for the end user, except:

a. The warranty period for the battery pack included with the Product is one (1) year from the date of shipment to the original purchaser.

b. The warranty period for accessories, replacement parts, and disposables including, but not limited to: filters, fuses, cannulas, tubing, shoulder bag and strap, accessory bag, AC charger, and DC charger is 90 days from the date of shipment to the original purchaser.

What this warranty does not cover: This warranty does not apply to any software included with the Product as the software warranty is included in the software license. This warranty does not cover damage or injury whether to the Products, personal property, or persons caused by accident, misuse, abuse, Acts of God, water ingress, unapproved ozone cleaning and disinfection methods, other unapproved cleaning and disinfection methods, repair or alteration by anyone other than Philips Respironics or its authorized service center, failure to operate in accordance with the terms of the operating manual and instructions, lack of reasonable care, the discontinuance of a network (e.g. 2G, 3G, etc.) by a carrier (e.g. ATT, Verizon, etc.), or other defects not related to material or workmanship. This warranty is not transferable. If Philips Respironics finds that a Product returned for service or the issue raised is not covered under this limited warranty, Philips Respironics may charge an evaluation fee and return shipping.

What Philips Respironics will do: If a Product does not meet the warranty above in the first 90 days after the original shipment date, Philips Respironics will replace the device with a new Product. Thereafter, if a Product fails to conform to the warranties set forth above during the applicable warranty period, Philips Respironics will repair or replace the Product or refund the original purchase price, in Philips Respironics sole discretion. Philips Respironics may use new or remanufactured assemblies, components, and parts in repair and new or recertified refurbished devices for replacement. The balance of the original warranty period will apply to any Product or component of a Product repaired or replaced under this warranty.

Warranty Disclaimer; Limitation of Liability: EXCEPT AS SET FORTH IN THIS LIMITED WARRANTY, PHILIPS RESPIRONICS MAKES NO WARRANTIES, EXPRESSED OR IMPLIED, STATUTORY OR OTHERWISE, REGARDING THE PRODUCT OR ITS QUALITY OR PERFORMANCE. PHILIPS RESPIRONICS SPECIFICALLY DISCLAIMS THE IMPLIED WARRANTY OF MERCHANTABILITY AND THE IMPLIED WARRANTY OF FITNESS FOR A PARTICULAR PURPOSE. IN NO EVENT WILL PHILIPS RESPIRONICS MAXIMUM LIABILITY UNDER THESE WARRANTIES EXCEED THE ORIGINAL PURCHASE PRICE OR WILL PHILIPS RESPIRONICS BE LIABLE FOR ANY ECONOMIC LOSS, LOSS OF PROFITS, OVERHEAD, OR SPECIAL, INCIDENTAL, OR CONSEQUENTIAL DAMAGES. Repair, replacement, or return of purchase price by Philips Respironics is the original purchaser's sole and exclusive remedy under this warranty.

This warranty gives you specific legal rights, and you may also have other rights that vary from state to state. Some states do not allow the exclusion or limitation of incidental or consequential damages, so the above exclusion and limitations may not apply to you.

How to get warranty support: Patients, contact your local authorized Philips Respironics dealer. Dealers, contact Respironics, Inc. at:

1001 Murry Ridge Lane

Murrysville, Pennsylvania 15668-8550

+1-724-387-4000

Note: For Australian and New Zealand customers this warranty replaces the warranty included with the user manual.

1. The following statement is provided to a customer who is a consumer under the Australian Consumer Law: Our goods come with guarantees that cannot be excluded under the Australian Consumer Law. You are entitled to a replacement or refund for a major failure and for compensation for any other reasonably foreseeable loss or damage. You are also entitled to have the goods repaired or replaced if the goods fail to be of acceptable quality and the failure does not amount to a major failure. This guarantee applies in addition to other rights and remedies available to the consumer under law.

2. The following statement is provided to a customer who is a consumer under the Consumer Guarantees Act 1993 (NZ): For consumers under the Consumer Guarantees Act 1993 (NZ) who purchase the goods for personal, domestic or household use: Our goods come with guarantees that cannot be excluded under the Consumer Guarantees Act 1993 (NZ).

3. Philips warrants that the products shall be free from defects of workmanship and materials, and will perform in accordance with the product specifications for a period of two (2) years from the date of purchase from an authorised Philips Homecare Provider. This Warranty covers the replacement or repair at the option of Philips, of any product that has a manufacturing or material defect that is not the result of normal wear and tear, or a natural characteristic of the material used. This Warranty is not transferable and does not cover products used for commercial purposes, and it does not apply to any consumable items (including but not limited to filters, masks, tubes and humidifier chambers).

4. The customer is responsible for returning the product to an authorised Philips Homecare Provider, and for collecting the product from the Homecare Provider after repair or replacement, at its own cost. Philips is responsible only for the freight cost of transporting the product between the Homecare Provider and Philips. Philips reserves the right to charge an evaluation and postage fee for any returned product where no problem is found following evaluation.

5. This Warranty does not cover:

- products purchased outside of Australia and New Zealand;
- any damage caused as a result of misuse or abuse, modification, tampering with or alteration of the product, pest infestation, liquid egress into the product, or unapproved ozone cleaning and disinfection methods;
- contamination due to cigarette, pipe, cigar or other smoke;
- failure to follow manufacturer's instruction for use as per the user manual;
- defects that are a consequence of repairs to a product made or attempted by a service provider other than one approved by Philips;
- products that have been subjected to incorrect electrical supply or inconsistent electrical supply or used with inappropriate accessories.

6. This Warranty is not transferrable in the event of any resale or transfer of products.

7. To make a claim under this Warranty, contact your Homecare Provider or Philips:

AUSTRALIA

Philips Electronics Australia Limited

65 Epping Road, North Ryde

NSW 2113 Australia

Tel: 1300 766 488

Email: repairs-src@philips-easyconnect.com

NEW ZEALAND

Philips New Zealand Commercial Ltd
Level 3, 123 Carlton Gore Road
New Market Auckland 1023 New Zealand
Tel: 0800 251 400
Email: repairs-src@philips-easyconnect.com

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