



Trilogy 100, Trilogy 200,
Trilogy 0₂ & Trilogy 202
Service & Technical Information



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REVISION HISTORY

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CHAPTER 1: INTRODUCTION

1.0 CHAPTER OVERVIEW

This chapter provides an introduction for the Trilogy Ventilator as well as contact and service training information.

1.1 TRILOGY 100 & TRILOGY 200 VENTILATORS INTENDED USE

The Trilogy 100 & Trilogy 200 systems provide continuous or intermittent ventilatory support for the care of individuals who require mechanical ventilation. The Trilogy 100 & Trilogy 200 are intended for pediatric through adult patients weighing at least 5 kg (11 lbs).

The device is intended to be used in home, institution/hospital, and portable applications such as wheelchairs and gurneys, and may be used for both invasive and non-invasive ventilation. It is not intended to be used as a transport ventilator.

The system is recommended to be used only with various combinations of Respironics-approved patient circuit accessories, such as patient interface devices, humidifiers, water traps, and circuit tubing.

1.2 TRILOGY O₂ & TRILOGY 202 INTENDED USE

The Respironics Trilogy O_2 & Trilogy 202 systems provides continuous or intermittent ventilatory support for the care of individuals who require mechanical ventilation with or without FiO_2 blending. Trilogy O_2 & Trilogy 202 are intended for pediatric through adult patients weighing at least 5 kg (11 lbs.).

The device is intended to be used in hospitals and institutions, and for portable applications such as wheelchairs and gurneys only when in an institutional setting. It may be used for both invasive and non-invasive ventilation. It is not intended to be used as a transport ventilator.

The system is recommended to be used only with various combinations of Respironics-approved patient circuit accessories, such as patient interface devices, humidifiers, water traps, and circuit tubing.

1.3 TRILOGY 100 & TRILOGY 200 SYSTEM OVERVIEW

This ventilator provides both pressure control and volume modes of therapy. The device can provide non-invasive or invasive ventilation. It can be used to provide total therapy to patients as they progress from non-invasive to invasive ventilation.

When prescribed, the device provides numerous special features to help make patient therapy more comfortable. For example, the ramp function allows you to lower the pressure when trying to fall asleep. The air pressure will gradually increase until the prescription pressure is reached. Additionally, the Flex comfort feature provides increased pressure relief during the expiratory phase of breathing.

The ventilator can be operated using several different power sources, including an internal Lithium-Ion battery. This battery is automatically used when the detachable Lithium-Ion battery pack, external Lead Acid battery, or AC power are not available.

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1.4 TRILOGY O₂ & TRILOGY 202 SYSTEM OVERVIEW

This ventilator provides both pressure control and volume modes of therapy. The device can provide non-invasive or invasive ventilation. It can be used to provide total therapy to patients as they progress from non-invasive to invasive ventilation.

When prescribed, the device provides numerous special features to help make therapy more comfortable. For example, the ramp function allows you to lower the pressure when trying to fall asleep. The air pressure will gradually increase until the prescription pressure is reached. Additionally, the Flex comfort feature provides increased pressure relief during expiratory phase of breathing.

The ventilator can be operated using several different power sources, including an internal Lithium-Ion battery. This battery is automatically used when the detachable Lithium-Ion battery pack, external Lead Acid battery, or AC power are not available.

This ventilator is equipped with an oxygen blending module which allows oxygen to be delivered to the patient within a range of 21% to 100% concentration.

1.5 SERVICE TRAINING

Respironics offers service training for the Trilogy devices. Training includes complete disassembly of the device, troubleshooting subassemblies and components, setup of test equipment, and necessary testing. For more information, contact the Service Marketing department at:

E-mail: service.operations@respironics.com Phone: (724) 755-8220 Fax: (724) 755-8230

1.6 Service/Technical Support Statement

For technical assistance, please contact Respironics Customer Satisfaction.

U.S.A. and Canada Phone:1-800-345-6443 Fax: 1-800-886-0245

International Phone: 1-724-387-4000

Fax: 1-724-387-5012



CHAPTER 2: WARNINGS, CAUTIONS, & NOTES

2.0 CHAPTER OVERVIEW

Warnings, cautions, and notes are used throughout this manual to identify possible safety hazards, conditions that may result in equipment or property damage, and important information that must be considered when performing service and testing procedures. Please read this chapter carefully before servicing Trilogy Ventilators.

WARNING

Warnings indicate the possibility of injury to people.

CAUTION

Cautions indicate the possibility of damage to the device.

NOTE

Notes are used to emphasize a characteristic or important consideration.

2.1 WARNINGS

WARNINGS

- Electronic components used in this device are subject to damage from static electricity. Repairs made to this device must be performed only in an antistatic, Electro-static Discharge (ESD)-protected environment.
- To assure the safety of the service technician and the specified performance of the device, Respironics requires that only trained technicians perform any repairs or adjustment to the Trilogy devices.
- High voltages are present inside this device. To avoid electrical shock, disconnect the electrical supply before attempting any repairs on the device.
- DO NOT immerse this device into any fluids.



2.2 CAUTIONS

CAUTIONS

- Federal law (US) restricts this device to sale by, or on the order of, a physician.
- Care should be taken to avoid exposure of Trilogy Ventilators to temperatures at or near the extremes of those specified in the Specifications Chapter of this manual. If exposure to such temperatures has occurred, the device should be allowed to return to room temperature before being turned on.
- Never place liquids on or near Trilogy Ventilators.
- To avoid electrical shock, disconnect the electrical supply before cleaning Trilogy Ventilators.
- The information in this manual is provided for service personnel reference.

2.3 Notes

NOTES

- Additional Warnings, Cautions and Notes are located throughout this manual.
- Refer to the Trilogy Provider Manuals for additional Warnings, Cautions and Notes.

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CHAPTER 3: SPECIFICATIONS, CLASSIFICATIONS, & SYSTEM FEATURES

3.0 CHAPTER OVERVIEW

This chapter identifies the specifications, classifications, & system features for the Trilogy Ventilators.

3.1 Environmental Specifications

	Operating	Storage
Temperature	5° C to 40° C	-20° C to 60° C
Relative Humidity	15 to 95% (non-condensing)	15 to 95% (non-condensing)
Atmospheric Pressure	110 kPa to 60 kPa	N/A

The operating range for the charging of the internal and detachable batteries is 10° C to 30° C. The internal and detachable batteries will power the ventilator for the full operating range of 5° C to 40° C.

Accuracies stated in this manual are based on specific environmental conditions. For stated accuracy, the environmental conditions are: Temperature: 20-30° C, Humidity: 50% relative, Altitude: nominally 380 meters

3.2 TRILOGY 100 & TRILOGY 200 PHYSICAL SPECIFICATIONS

Dimensions	16.68 cm L x 28.45 cm W x 23.52 cm H
Weight	Approximately 5 kg (with the detachable battery installed)

3.3 TRILOGY O₂ & TRILOGY 202 PHYSICAL SPECIFICATIONS

Dimensions	21.13 cm L x 28.45 cm W x 23.52 cm H
Weight	Approximately 6.1 kg (with the detachable battery installed)



3.4 ELECTRICAL SPECIFICATIONS

AC Voltage Source	100 to 240 VAC, 50/60 Hz, 2.1 A
Detachable Battery:	Voltage: 14.4 VDC Capacity: 71 Wh Chemistry type: Lithium-Ion
Internal Battery:	Voltage: 14.4 VDC Capacity: 71 Wh Chemistry type: Lithium-Ion
Type of Protection Against Electric Shock:	Class II/Internally Powered Equipment
Degree of Protection Against Electric Shock:	Type BF Applied Part
Degree of Protection Against Ingress of Water:	Device: Drip Proof, IPX1
Mode of Operation:	Continuous
Fuses:	There are no user-replaceable fuses.
Power Consumption:	2.1 A @ 100 VAC = 210 Watts of Power Consumption

3.5 PRESSURE SPECIFICATIONS

Output:	4 to 50 pressure units (may be cmH ₂ O, hPA, or
	mBar depending on device setup)



3.6 SD CARD & SD CARD READER

Use only SD cards and SD card readers available from Respironics or the following:

- SanDisk[®] Standard SD Card 2.0 GB Respironics Part Number: 1053952
- SD Card Reader/Writer Respironics Part Number: 1047300

3.7 STANDARDS COMPLIANCE

This device is designed to conform to the following standards:

- IEC 60601-1: Medical electrical equipment Part 1: General requirements for safety
- IEC 60601-1-2: General requirements for safety Collateral standard: Electromagnetic compatibility Requirements and tests
- ISO 10651-2-Lung ventilators for medical use -- Particular requirements for basic safety and essential performance -- Part 2: Home care ventilators for ventilator-dependant patients
- RTCA-D0160F section 21, category M; Emission of Radio Frequency Energy (Trilogy 200 Only)



3.8 CONTROL ACCURACY

PARAMETER	RANGE	ACCURACY	
IPAP	4 to 50 ¹ pressure units ²	Greater of 2 pressure units or 8% of setting	
EPAP	0 to 25 pressure units for Active Circuits 4 to 25 pressure units for Passive Circuits	Greater of 2 pressure units or 8% of setting	
CPAP	4 to 20 pressure units	Greater of 2 pressure units or 8% of setting	
PEEP	0 to 25 pressure units for Active Circuits 4 to 25 pressure units for Passive Circuits	Greater of 2 pressure units or 8% of setting ⁴	
Pressure	4 to 50 pressure units	Greater of 2 pressure units or 8% of setting	
Pressure support	0 to 30 pressure units ³	Greater of 2 pressure units or 8% of setting ⁴	
Tidal Volume	50 to 2000 ml ⁵	Greater of 10 ml or 10% of setting (Active Circuits) Greater of 15 ml or 15% of setting (Passive Circuits)	
Breath Rate	0 to 60 BPM for AC mode 1 to 60 BPM for all other modes	Greater of ± 1 BPM or ±10% of the setting	
Timed Inspiration	0.3 to 5.0 seconds	± .1 second	
Rise Time	1 to 6 ⁶	± 2 pressure units (the device will increase to a pressure of .67* (IPAP - EPAP) ± 2 pressure units @ the set rise time multiplied by 100 ms for pressure supports less than or equal to 25.)	
Ramp Start Pressure	0 to 25 pressure units for Active Circuits 4 to 25 pressure units for Passive Circuits 4 to 19 pressure units in CPAP mode	8% of setting + 2% Full Scale	
Ramp Length	Off, 5 to 45 minutes	<u>+</u> 2 minutes	
Flex	Off, 1 to 3 ⁷	N/A	
Flow Trigger Sensitivity	1 to 9 l/min	N/A	
Trilogy 100, Trilogy 200, & Trilogy 202, Flow Cycle	10 to 90%	N/A	
Trilogy O ₂ Flow Cycle	10 to 40%	N/A	
Apnea Rate	4 to 60 BPM	Greater of 1 BPM or 10% of setting	
FiO2 Output (Trilogy O ₂ & Trilogy 202 Only)	21% to 100%	21% to 50% is ±3% 50% to 95% is ±5% 100% is -5%	
O2 Input Pressure Rating (Trilogy O ₂ & Trilogy 202 Only)	40 to 87 PSI		

Specifications listed above are based on using a standard circuit (1.8 meter tubing - REF 622038; Passive Exhalation Device - REF 1040417; Active Exhalation Device with PAP - REF 1053716)

¹Limited to 25 pressure units when using the Bi-Flex feature in S mode.

²Pressure units may be cmH₂O, hPa, or mBar depending on device setup.

³The difference between the Inspiratory Pressure and the Expiratory Pressure must never be more than 30 pressure units.

⁴Pressure Support and PEEP not to exceed 50 pressure units.

⁵Reflects compensation for BTPS.

⁶The range of values correspond to the tenths of seconds (e.g., a setting of 4 indicates a Rise Time of 0.4 seconds).

⁷Flex is not available when AVAPS is active. Flex is not available with Active Circuits.



3.9 MEASURED PATIENT PARAMETERS

PARAMETER	RANGE	ACCURACY	
Vte/Vti	0 to 2000 ml	Greater of 15ml or 15% of reading	
Minute Ventilation	0 to 99 l/min	Calculation based on measured Vte or Vti and Respiratory Rate	
Estimated Leak Rate	0 to 200 l/min	N/A	
Respiratory Rate	0 to 80 BPM	Greater of 1 BPM or 10% of reading	
Peak Inspiratory Flow	0 to 200 l/min	3 l/min plus 15% of reading	
Peak Inspiratory Pressure	0 to 99 pressure units	Greater than 2 pressure units or 10% of reading	
Mean Airway Pressure	0 to 99 pressure units	Greater than 2 pressure units or 10% of reading	
% Patient Triggered Breaths	0 to 100%	N/A	
I:E Ratio	9.9- 1: 1-9.9	Calculation based on Inspiratory Time and Expiratory Time	
All flows and volumes are meas	cured at BTPS conditions.	1	

3.10 TRILOGY 100 SPONTANEOUS BREATHING DURING POWER FAILURE CONDITIONS

FLOW SET POINT (LPM)		RESISTANCE RE UNITS)	EXPIRATORY RESISTANCE (PRESSURE UNITS)	
	Active Circuit	Passive Circuit	Active Circuit	Passive Circuit
30	< 2.0	< 1.0	< 1.5	< 1.2
60	< 10.0	< 4.0	< 4.0	< 3.7

3.11 TRILOGY 200, TRILOGY O₂, & TRILOGY 202 SPONTANEOUS BREATHING DUR-ING POWER FAILURE CONDITIONS

FLOW SET POINT	INSPIRATORY RESISTANCE (PRESSURE UNITS)			EXPIRATORY RESISTANCE (PRESSURE UNITS)		
(LPM)	Active Circuit	Active Circuit with Proximal Flow Sensor	Passive Circuit	Active Circuit	Active Circuit with Proximal Flow Sensor	Passive Circuit
30	< 3.0	< 3.5	< 2.0	< 2.0	< 2.0	< 1.5
60	< 9.0	< 10.5	< 5.0	< 3.0	< 4.5	< 4.0

3.12 RESISTANCE VALUES

During a ventilator failure, both the inhalation resistance and the exhalation resistance, measured at the patient connection port, will not exceed 10 pressure units at 60 l/min. or 0.8 pressure units at 5 l/min.



3.13 WEEE/ROHS RECYCLING DIRECTIVES

If you are subject to the WEEE/RoHS recycling directives, refer to www.respironics.com for the passport for recycling this product and the batteries.



3.14 EMC INFORMATION

3.14.1 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
Harmonic emissions IEC 61000-3-2	Class A	establishments and those directly connected to the public low-voltage power supply network that supplies building used for
Voltage fluctuations/Flicker emissions IEC 61000-3-3	Complies	domestic purpose.



3.14.2 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	±6 kV contact ±8 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 for supply mains ±1 kV for input/output lines	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 ±2 kV for common mode	Mains power quality should be that of a typical home or hospital environment.	
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 40% U_T $(60\%$ dip in U_T) for 5 cycles 70% U_T (30% dip in U_T) for 25 cycles $<5\%$ U_T (>95% dip in U_T) for 5 sec	$<5\%$ U_T $(>95\%$ dip in U_T) for 0.5 cycle 40% U_T $(60\%$ dip in U_T) for 5 cycles 70% U_T (30% dip in U_T) for 25 cycles $<5\%$ U_T (>95% dip in U_T) for 5 sec	Mains power quality should be that of a typical home 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 supply or a battery.	
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	3 A/m	3 A/m	Power frequency magnetic fields should be at levels characteristic of a typical home or hospital environment.	
NOTE: U_T is the a.c. mains voltage prior to application of the test level.				

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3.14.3 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 (FDA)	ELECTROMAGNETIC ENVIRONMENT - GUIDANCE
Conducted RF IEC 61000-4-6	3 Vrms 150 kHz to 80 MHz outside ISM bands ^a	3 V	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. Recommended separation distance: $d = 1.2 \ \sqrt{P}$
	10Vrms 150 kHz to 80 MHz in ISM bands ^a	10 V	$d = 1.2 \sqrt{P}$
Radiated RF IEC 61000-4-3	10 V/m 80 MHz to 2.5 GHz	10 V/m 26 MHz to 2.5 Gz	$d = 1.2 \ \sqrt{P} \ 80 \ \text{MHz} \text{ to } 800 \ \text{MHz}$ $d = 2.3 \ \sqrt{P} \ 800 \ \text{MHz} \text{ to } 2.5 \ \text{GHz}$
			where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in meters (m).
			Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, should be less than the compliance level in each frequency range.b
			Interference may occur in the vicinity of equipment marked with the following symbol:
			((·*))

NOTE 1 At 80 MHz and 800 MHz, the higher frequency range applies.

NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects, and people.

a Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the device is used exceeds the applicable RF compliance level above, the device should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the device.

b Over the frequency range 150 kHz to 80 MHz, the field strengths should be less than 3 V/m.



3.14.4 RECOMMENDED SEPARATION DISTANCES BETWEEN PORTABLE AND MOBILE RF COMMUNICATIONS EQUIPMENT

The device is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of this device can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and this device as recommended below, according to the maximum output power of the communications equipment.

RATED MAXIMUM POWER OUTPUT OF TRANSMITTER (WATTS)	SEPARATION DISTANCE ACCORDING TO FREQUENCY OF TRANSMITTER (METERS)			
	150 kHz to 80 MHz outside ISM Bands $d = 1.2 \sqrt{P}$	150 kHz to 80 MHz outside ISM Bands $d = 1.2 \sqrt{P}$	80 MHz to 800 MHz $d = 1.2 \sqrt{P}$	800 MHz to 2.5 GHz $d = 2.3 \sqrt{P}$
0.01	0.12	0.12	0.12	0.23
0.1	0.38	0.38	0.38	0.73
1	1.2	1.2	1.2	2.3
10	3.8	3.8	3.8	7.3
100	12	12	12	23

For transmitters rated at a maximum output power not listed above, the recommended separation distance **d** in meters (m) can be estimated using the equation applicable to the frequency of the transmitter, where P is the maximum output power rating of the transmitter manufacturer.

Note 1: At 80 MHz and 800 MHz, the higher frequency range applies.

Note 2: The ISM (industrial, scientific and medical) bands between 150 kHz and 80 MHz are 6.765 MHz to 6.795 MHz;13.553 MHz to 13.567 MHz; 26.957 MHz to 27.283 MHz; and 40.66 MHz to 40.70 MHz.

Note 3: An additional factor of 10/3 is used in calculating the recommended separation distance for transmitters in the ISM frequency bands between 150 kHz and 80 MHz and in the frequency range of 80 MHz and 2.5 GHz to decrease the likelihood that mobile/portable communications equipment could interference if it is inadvertently brought into patient areas.

Note 4: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects, and people.

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3.15 FRONT PANEL FEATURES

The front panel contains the control buttons, visual indicators, and display screen.

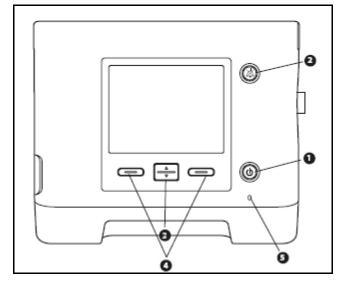


FIGURE 3-1: FRONT PANEL CONTROLS & DISPLAY SCREEN

NOTE

When you start therapy, the display backlight and the backlights on the buttons turn on, the red and yellow alarm LEDs turn on momentarily, and an audible indicator sounds to indicate that therapy has started. The Startup screen appears on the display.

WARNING

To make sure the device is operating properly at start-up, always verify that the audible tone sounds and the Audio Pause LED lights red and then yellow momentarily.

3.15.1 Buttons

The following buttons are included on the front panel of the device.

- Start/Stop Button This button turns the airflow on or off, starting or stopping therapy.
- 2. **Alarm Indicator and Audio Pause Button** This button serves two purposes: it temporarily silences the audible portion of an alarm, and it also acts as an alarm indicator. When silencing an alarm, if the cause of the alarm is not corrected, the alarm sounds again after one minute. Each time the button is pressed, the alarm silence period resets to one minute.
- 3. **Up/Down Button** This button allows you to navigate the display menu and edit device settings.
- 4. **Left and Right Buttons** These buttons allow you to select display options or perform certain actions specified on-screen.



3.15.2 VISUAL INDICATORS

Several power and alarm indicators appear on the front panel.

- AC Power LED In the lower right hand corner of the front panel, a green LED indicates that the AC power is applied to the device. This light remains on as long as adequate AC power is available.
- 2. **Keypad Backlight LEDs** The Start/Stop, Up/Down, and Left/Right buttons all have a white LED that lights up if the keypad backlight is turned on in the device Options menu.
- 3. Red Alarm LED On the Audio Pause button, a red light flashes to indicate a high priority alarm.
- 4. **Yellow Alarm LED** On the Alarm Indicator/Audio Pause button, a yellow light flashes to indicate a medium priority alarm. A solid yellow light indicates a low priority alarm.

3.15.3 DISPLAY SCREEN

The display screen allows you to view settings, system status information, real-time patient data, alarms, and logs. You can also modify certain settings on the display screen.

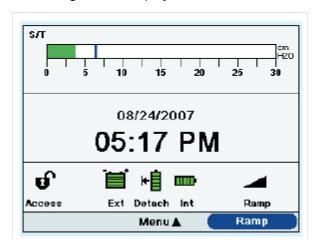


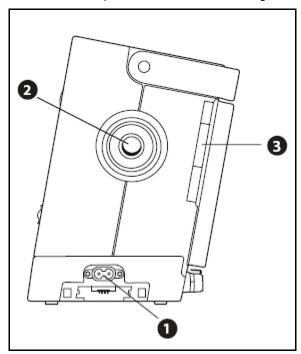
FIGURE 3-2: SAMPLE DISPLAY SCREEN

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3.16 TRILOGY 100 & TRILOGY 200 SIDE PANEL FEATURES

The ventilator's side panels contain the following connectors and features.



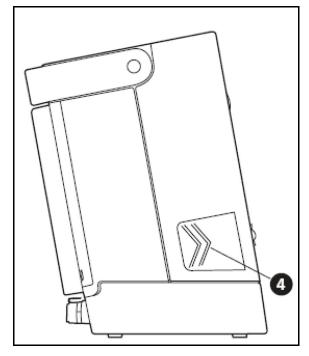


FIGURE 3-3: RIGHT & LEFT SIDE PANEL

- 1. **AC Power Inlet** You can plug the AC power cord into this connector, located on the right side of the ventilator.
- 2. **Breathing Circuit Connection** The breathing circuit connector is located on the right side of the device. You can connect your circuit tubing system here.
- 3. **Exhalation Porting Block** The porting block used here depends on whether you are using the Whisper Swivel II or the active exhalation device. The Passive Exhalation Porting Block is shown here.
- Secure Digital (SD) Data Card Slot On the left side of the device is a slot for the optional SD
 Data Card. You can have the patient record usage and therapy information from the device on the
 SD card.



3.17 TRILOGY 100 & TRILOGY 200 REAR PANEL FEATURES

The ventilator's rear panel has the connectors and features.

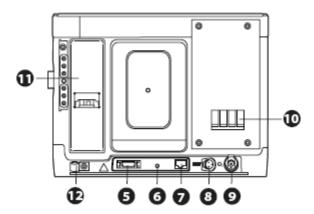


FIGURE 3-4: REAR PANEL

- 5. **Serial Connector** You can use this connector to connect the device to a computer running PC Direct or Alice Sleepware software or to other Respironics' devices such as Ailce 5 or AOM. Use the Trilogy RS232 Serial cable to connect the Trilogy to the external device or computer.
- 6. **Remote Alarm/Nurse Call Connector** If you are using an optional remote alarm or nurse call with the ventilator, you can connect the Respironics remote alarm adapter cable or nurse call adaptor cable to this connector.
- 7. **Ethernet Connector (when available)** You can connect a PC or router to this connector to upload therapy information to a secure website so you can review therapy information remotely or remotely troubleshoot and service the device.
- 8. **External Battery Connector (DC Power Inlet)** You can connect an external, stand-alone lead acid battery here, using the Respironics External Battery cable.
- 9. Oxygen (O₂) Inlet Connector If using low flow, supplemental oxygen, connect the oxygen source to this connector using one of the O₂ Inlet Quick Connects provided with the device.
- Air Inlet and Filter Insert the filter supplied with the device into the air inlet.
- 11. **Detachable Battery Pack Slot** If you are using the Respironics Lithium-lon detachable battery pack to power the device, attach it here.
- 12. **Cord Retainer** Wrap the power cord around this cord retainer to prevent someone from accidentally disconnecting the power cord.

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3.18 TRILOGY O_2 & TRILOGY 202 SIDE PANEL FEATURES

The ventilator's side panels contain the following connectors and features.

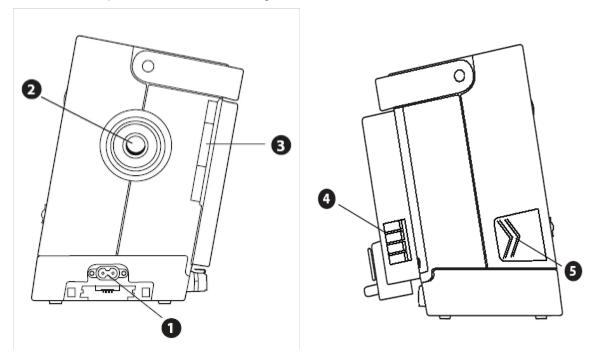


FIGURE S: RIGHT & LEFT SIDE PANEL

- 1. **AC Power Inlet** You can plug the AC power cord into this connector, located on the right side of the Trilogy Ventilator.
- 2. **Breathing Circuit Connection** The breathing circuit connector is located on the right side of the device. You can connect your circuit tubing system here.
- 3. **Exhalation Porting Block** the porting block used here depends on whether you are using the passive exhalation device or the active exhalation valve. The Passive Exhalation Porting Block is shown here. If you are using the active exhalation device, attach the Active Exhalation Porting block.
- 4. Air Inlet and Filter Insert the filter supplied with the device into the air inlet.
- 5. **Secure Digital (SD) Data Card Slot -** On the left side of the device is a slot for the optional SD Data Card. You can have the patient record usage and therapy information from the device on the SD card.



3.19 TRILOGY O₂ & TRILOGY 202 REAR PANEL FEATURES

The ventilator's rear panel has the connectors and features.

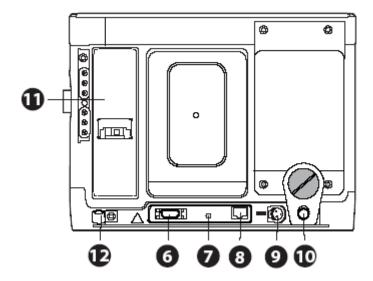


FIGURE A: REAR PANEL

- **6. Serial Connector** You can use this connector to connect the device to a computer running PC Direct or Alice Sleepware software or to other Respironics' devices such as Ailce 5 or AOM.
- 7. **Remote Alarm Connector** If you are using an optional remote alarm with the Trilogy, you can connect the Respironics remote alarm adapter cable or nurse call cable to this connector.
- **8. Ethernet Connector** You can connect a PC or router to this connector to upload therapy information to a secure website so you can review therapy information remotely or remotely trouble-shoot and service the device.
- **9. External Battery Connector (DC Power Inlet)** You can connect an external, stand-alone lead acid battery here, using the Respironics External Battery cable.
- Oxygen (O₂) Blending Module If using high pressure supplemental oxygen, connect the oxygen source to this DISS connector.
- 11. **Detachable Battery Pack Slot** If you are using the Respironics Lithium-Ion detachable battery pack to power the device, attach it here.
- **12. Cord Retainer** Wrap the power cord around this cord retainer to prevent someone from accidentally disconnecting the power cord.

3.20 Power Sources

The ventilator accesses power from potential sources in the following order:

- AC Power
- External Battery
- Detachable Battery Pack

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Internal Battery

WARNING

Before connecting the battery, shutoff and disconnect any oxygen sources connected to the ventilator, and move the battery at least 1.83 meters (6 feet) away from any oxygen sources.

3.20.1 AC POWER SOURCE

An AC power cord is provided with the device. You can connect the pronged end of the power cord to a wall outlet that is not controlled by a wall switch and connect the socket end of the cord to the power inlet on the back of the device.

3.20.2 EXTERNAL BATTERY

CAUTION

- Do not use the same external battery to operate both the ventilator and any other equipment.
- An external battery should only be connected to the ventilator using the Respironics Trilogy
 External Battery Cable. This cable is fused, pre-wired and properly terminated to ensure
 safe connection to a standard deep cycle lead acid battery. Use of any other adaptor or
 cable may cause improper operation of the ventilator.

The ventilator can operate from a 12 VDC deep cycle marine-type (lead acid) battery using the Respironics Trilogy External Battery Cable. This cable is pre-wired and properly terminated to ensure safe connection of an external battery to the ventilator. Battery operating time depends on the characteristics of the battery and usage of the device.

Due to a variety of factors, including battery chemistry, battery age, and use profile, the capacity of the external battery as shown on the Trilogy display is only an estimate of the actual remaining capacity.

Refer to the instructions supplied with the Respironics External Battery Cable for detailed information on how to operate the device using an external battery.



3.20.3 DETACHABLE BATTERY

CAUTION

- The detachable and internal batteries wear out based on the amount of use (hours or full charge-discharge cycles). The battery capacity and life are also reduced by operation at higher temperatures.
- Only use the Respironics Trilogy Detachable Battery with the device.
- Prolonged operation or storage at elevated temperatures may reduce the service life of the detachable or internal battery and other internal components of the ventilator.

Respironics offers a detachable Lithium-ion battery pack. To use the detachable battery pack, snap the battery into place on the back of the ventilator. When the device is not connected to an AC power source or an external battery, the detachable battery will power the device, if attached. The length of time the ventilator will operate on battery power depends on many factors such as device settings, battery charge level, and condition or age of the battery. When fully charged, a new battery can power the ventilator for approximately three hours under typical patient conditions.

Whenever the ventilator is connected to AC power, it will automatically recharge the detachable battery pack. A completely discharged detachable battery will reach 80% charge status within 8 hours, when charging at approximately 23° C ambient temperatures.

One side of the detachable battery has a set of LEDs that indicate the amount of charge left on the battery. You can press the button below the LEDs to view how much charge remains:

LED	BATTERY CAPACITY
All 5 LEDs are lit	80-100% capacity
4 LEDs are lit	60-79% capacity
3 LEDs are lit	40-59% capacity
2 LEDs are lit	20-39% capacity
1 LED is lit	10 to 19% capacity
1 LED flashes	1 to 9% capacity
0 LEDs are lit	0% capacity

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3.20.4 INTERNAL BATTERY

CAUTION

The internal battery is NOT intended to serve as a primary power source. It should only be used when other sources are not available, or briefly when necessary; for example when changing power sources.

The device contains an internal battery that can be used as a backup power source. It is intended for use during short periods while switching between external power sources, emergency situations, or short durations when the user needs to be mobile. The length of time the ventilator will operate on internal power depends on many factors such as device settings, battery charge level, and condition or age of the battery. When fully charged, a new battery can power the ventilator for approximately three hours under typical patient conditions.

Whenever the ventilator is connected to AC power, it will automatically recharge the internal battery. A completely discharged internal battery will reach 80% charge status within 8 hours, when charging at 23° C ambient temperature.



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CHAPTER 4: THEORY OF OPERATION

4.0 CHAPTER OVERVIEW

This document describes the theory of operation for the seven printed circuit boards employed by the Trilogy ventilator support system.

4.1 AC/DC POWER SUPPLY

The AC/DC power supply is capable of accepting universal AC input supply of 100 - 240VAC @ 50/60Hz. It generates a 30±2VDC nominal output with ripple voltage no greater than 250mVp-p over its entire operating range. It is capable of delivering 3A (90 Watts) continuous and 5A (150Watts) peak (10% duty cycle) output current.

The AC/DC power supply uses isolated fly-back topology. It incorporates the "TOPSwitch" (U1) from Power Integrations with an internal high voltage power MOSFET. It operates at a switching frequency of 66±7 KHz. The output of the power supply is regulated via a 3-terminal precision voltage regulator (U2).

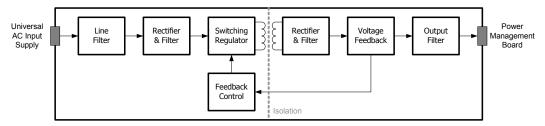


FIGURE 4-1 AC/DC POWER SUPPLY BLOCK DIAGRAM

The DC output of the AC/DC power supply is isolated from the AC input using the transformer (T1) and optoisolator (U3). The power supply can maintain 4000VAC isolation between primary and secondary, with less than 1mA leakage current when held for 10 seconds. The physical spacing and galvanic isolation maintained at 4mm. Slots are cut on the power supply board to satisfy adequate spacing where needed and to facilitate air flow.

The power supply is able to survive a DC short indefinitely and recover to normal operation after the DC short is removed. The operator should be careful and avoid touching any components on the power supply board while the unit is plugged into an AC source or even several minutes after AC power is removed. There is a risk of electrical shock that could cause serious injury or death. Assure C2 is thoroughly discharged before attempting to service any components.

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4.2 POWER MANAGEMENT BOARD

The Trilogy ventilator is designed to accept four power sources each providing enough power to support the complete functionality and performance of the ventilator. The power management board controls all these power sources for the Trilogy ventilator. The power sources are used according to the predefined hierarchy and rules set forth. The power management board is also capable of removing all power from the ventilator except for those devices powered by a low current supply to conserve power when the ventilator is turned off or running on battery. Lastly, the power management board provides a source of power diagnostic information through analog sense voltages and the SMBus to the CPU.

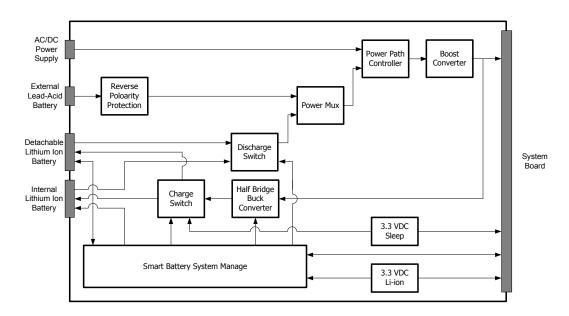


FIGURE 4-2 POWER MANAGEMENT BOARD BLOCK DIAGRAM

AC/DC power supply - It is the primary source of power for the Trilogy ventilator. It supplies 29±1VDC to the power management board and the Trilogy ventilator may draw up to a maximum of 3A continuous and 5A peak current from this power source. When this supply is available no current is drawn from any other remaining power source.

External Pb-Acid Battery - The Trilogy ventilator can operate on an external Pb-acid battery with output voltage between 11VDC to 28VDC when the AC/DC power supply is not available. The external Pb-acid battery input is reverse polarity protected (Q1). The Trilogy ventilator can not charge the external Pb-acid battery and can not use this source to charge the lithium ion batteries.

Detachable Lithium Ion Battery - This power source is used by the Trilogy ventilator when AC/DC power supply and external Pb-acid battery power supply is not available. It provides 12VDC to 16.4VDC supply to the ventilator and is rated for 4.16Ah @ 14.4V. The Trilogy ventilator may draw a maximum of 7.5A continuous or 14A peak current from the detachable lithium ion battery source. This battery is charged by the AC/DC power supply only when its capacity is less than 100%, the environment is safe to charge the battery and when charging would not overburden the AC/DC power supply.

Internal Lithium Ion Battery - This is the fourth and last available power source for the Trilogy ventilator. It is permanently located inside the unit and is identical to the detachable lithium ion battery (i.e. 12VDC to 16.4VDC, 4.2 Ah capacity). If the internal lithium ion battery is totally discharged and no other power source is available, the Trilogy ventilator will turn OFF, enable the power fail alarm and enter a low power state.

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Boost Converter - The boost converter (U1) generates the regulated 29VDC supply used by the system board. When the AC/DC power supply output voltage is greater than 28VDC the boost converter is OFF. If the AC/DC power supply is not available one of the battery sources is selected. The boost converter will then boost that battery voltage to 29VDC. When the input to the boost converter falls below 7VDC it is disabled.

Power Path Controller - The power path controller (U6) selects either the battery power supply or the AC/DC power supply to run the ventilator. When AC/DC supply is available the power path controller turns ON the AC/DC supply MOSFET (Q20) allowing the boost converter to receive its power from the AC/DC supply. When AC/DC supply in not available the power path controller turns ON the two parallel battery MOSFETs (Q25 & Q26) allowing the boost converter to receive its power from the battery supply. Initially, upon power up the circuitry powers the system with these MOSFETs acting as forward biased diodes until the sense and status lines for the power path controller have reached equilibrium.

Power Mux - The power path switch driver (U2) selects either the Pb-acid battery source or the lithium ion battery source to power the system. It does so by controlling 4 N-channel MOSFETs (Q11, Q12, Q13, & Q14) configured as two back to back ideal diode sources. This circuitry also disables all battery current when Trilogy enters the low power state by turning the N-Channel MOSFETs OFF.

Smart Battery System Manager - The smart battery system manager (U3) controls both the internal and detachable lithium ion batteries. Both batteries are equipped with smart technology and communicate with the power management board using the standard SMBus protocol. The smart battery system manager allows the CPU to communicate with either battery and contains memory for battery status and charger information while simultaneously monitoring the status and alarms bits of both batteries to prevent hazardous operation of either battery.

The software queries battery information about once per second. If the detachable lithium ion battery is disconnected, the comparator (U3) will quickly switch power to the internal lithium ion battery. Thereby continuing to power the system before the software realizes that a disconnect has happened. This comparator also prevents the software from disabling the internal lithium ion battery when there is no detachable battery present. The detachable battery is only disabled by the software if the software determines it to be a counterfeit battery.

Each lithium ion battery will power the system until its capacity falls below 10% of its full charge capacity. When both the lithium ion batteries are at less than 10% of the full charge capacity, they share the current drawn by the ventilator. This is done such that the battery with the greater capacity provides more current and both batteries reach 0% capacity simultaneously.

When both lithium ion batteries are at less than 100% charge capacity, both batteries share a maximum of 2A charge current according to their relative charge state. This is done such that the battery with less capacity receives greater charge current and both batteries reach 100% capacity simultaneously. The charge voltage and current limit used for each lithium ion battery is set at 16.9V and 2A respectively using resistors (R39, R35). In addition, the current sense resistor (R26) disables the battery charging operation when the total current output from the power supply exceeds 4A to prevent overloading of the power supply.

The smart battery system manager also generates an internal 5VDC supply to control all internal functions from the greatest available power source. A voltage divider network (R28 & R38) sets the AC present threshold at 22VDC coming from the AC/DC power supply. The smart battery system manger (U3) uses an internal comparator with a 1.19V reference to determine if AC is present.

Half Bridge Buck Converter - The buck converter (Q17, Q18, L4, & C33) generates the lithium ion battery charge voltage for both the lithium ion batteries. The charge switch is controlled by the smart battery system manager (U3).

Charge Switch - The charge switches (Q3, Q4, Q5, & Q6) facilitate charge current sharing between the two lithium ion batteries when they simultaneously request a charge. It is controlled by the smart battery system manager (U3) which provides charge current inversely proportional to the relative battery capacity using the charge switch.

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Discharge Switch - The discharge switches (Q7, Q8, Q9 & Q10) are used to control the discharge of the lithium ion batteries. The smart battery system manager (U3) controls the discharging of the internal and detachable lithium ion battery. The input MOSFETs (Q7, Q8) are turned OFF to prevent any accidental charging of a battery if reverse current is measured in the battery discharge path.

3.3VDC Supply - There are two 3.3VDC supplies generated on the power management board. First, the linear voltage regulator (U12) generates a 3.3VDC supply used when the ventilator is in the low power mode. And second, the switching regulator (U9) generates a 3.3VDC supply used by the logic circuitry on the board. The switching regulator (U9) is in shutdown mode when the Trilogy ventilator is in low power mode.

4.3 SYSTEM BOARD

A system board is the central printed circuit board in the Trilogy ventilator. It contains circuitry to interface with all other printed circuit board assemblies in the Trilogy ventilator with the exception of the power supply board. It generates and manages most of the internal power supply voltages for the Trilogy ventilator and contains the complete motor control circuitry.

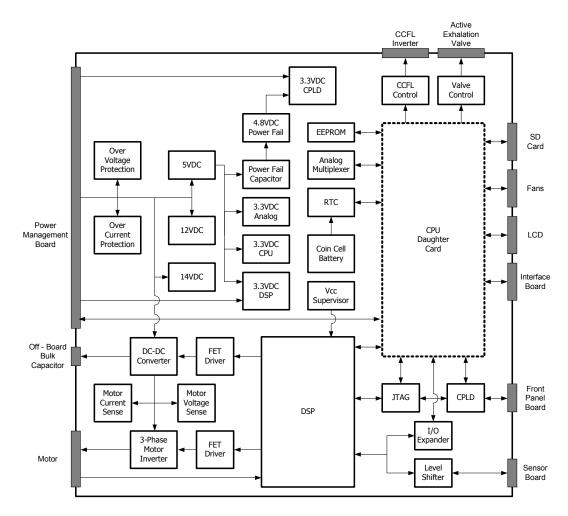


FIGURE 4-3 SYSTEM BOARD BLOCK DIAGRAM

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12VDC Supply - The dual switcher (U2) generates the 12VDC (2.5A max) supply using the 29VDC nominal supply from the power management. This supply is used by the MOSFET driver IC's, CCFL inverter, the front panel LEDs and to generate the 7VDC power supply for the sensor board.

5VDC Supply - The dual switcher (U2) generates the 5VDC (2.5A max) supply using the 29VDC nominal supply from the power management. This supply is used to generate the 3.3VDC supplies, power the motor hall sensors, power the system protection circuits and power some components on the interface board.

14VDC Supply - The step down switching regulator (U18) generates the 14VDC supply. This is a dedicated power supply to power the sensor board.

3.3VDC Supply - There are several 3.3VDC supplies generated on the system board.

- Buck regulator (U5) generates the 3.3VDC supply for the DSP. The 3.3VDC li-ion supply generated on the power management board acts as a back up to this supply
- Linear voltage regulator (U6) generates the 3.3VDC supply used by the analog signal conditioning circuits
- Step-down DC/DC converter (U4) generated the 3.3VDC supply for the CPU
- Linear voltage regulator (U3) generates the 3.3VDC supply for the CPLD

Power Fail Supply - The 5VDC supply is also used to charge the 2.5F ultra capacitor (C113). In the event of a power failure, the DC/DC converter (U24) uses the charge stored in the ultra capacitor to generate 4.8VDC nominal supply for power failure alarm generation.

Digital Signal Processor (DSP) - The Trilogy system board uses the DSP (U7) to control the motor and acts as a watchdog for the main CPU. It uses internal timers, interrupts, PWM, ADC and peripherals to run the motor control algorithm. It has six 12-bit ADC inputs available used to monitor motor voltage, DC to DC output voltage, motor current, system current, 1.2VDC reference voltage and motor temperature. It uses the SPI bus to communicate with the sensor board and the 16-bit I/O expander (U17).

Complex Programmable Logic Devices (CPLD) - The CPLD is used for power fail operations, low power mode, power up/wakeup and alarm functions of the Trilogy ventilator. It has a dedicated JTAG programming connector (J9) and operated of a 3.3VDC supply.

DC-DC Converter - The DC to DC converter controls the voltage and current applied to the motor to produce the desired speed. The converter consists of two N-channel FETs (Q3 & Q4) with an LC filter output stage. The LC filter reduces both the ripple current and ripple voltage. The FETs are driven by the half bridge power MOSFET driver (U10) controlled by the DSP. It also acts as a buffer between the motor control and the rest of the system through a blocking diode (CR10).

3 Phase Motor Inverter - The current through the six phases of the motor are controlled using the inverter. The inverter produces multi-phase current of variable frequency to provide the appropriate torque to control the motor speed. The inverter stages consist of three half bridge power MOSTFET drivers (U13, U15 & U16) controlling six N-channel MOSFETs (Q6 to Q11). The control signals for the driver are generated by the DSP.

Motor (external) - The motor used is a Brushless DC motor with low inductance, a WYE wound stator and 3 hall sensors located 120° apart (J5). A built in temperature sensor mounted on the motor board is used by the DSP to monitor the motor temperature.

Off Board Storage Capacitor - An off board bulk capacitor (J6) stores the regenerative energy of the motor during deceleration. It provides the ripple current necessary to accelerate and decelerate the motor and filter the motor free wheeling voltage from damaging the MOSFETs.

Over Voltage Protection - The comparator (U31) determines if an over voltage condition occurs on the high side of the DC-DC converter. If the measured voltage exceeds 35V, a precision current (U33 & Q28) sink is enabled to shunt current through the bleeder resistors (R174 & R175).

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Over Current Protection - The system current is measured by the high side current monitor (U12) in conjunction with the current sense resistor (R54). The buffer amplifiers (U36) drive the comparator (U29) with a current limit set at 5.5A with 500mA of hysteresis. When the system current increase above the current set point limit current to the motor is temporarily disabled and the motor spins freely until the current drops to a safe level.

Motor Current Sense - A current sense resistor (R39) measures the bidirectional current through three motor phases. The op-amp (U11: C) filters the current sense output before being used by the DSP to control the motor.

Motor Voltage Sense - The terminal voltage across the motor is monitored using a resistor divider network (R73 & R74) along with the op-amp circuit (U11:A). This signal is used by the DSP to control the motor.

Motor Temperature Sensing - Motor temperature monitored by the DSP using thermistor mounted on the motor. If motor temperature rises above the set point the DSP limits the current going to the motor to keep it cool.

Active Exhalation Valve Control - The active exhalation valve consists of a two valve system. One valve dumps the pilot pressure while the other provides proportional control of the pilot pressure. Both the valves are controlled by the CPU via connector on the system board (J10). To control the proportional valve the CPU generates an analog control voltage uses a 16-bit DAC (U27). This signal drives a precision current sink (U28 and Q5) controlling the current flow through the proportional active exhalation valve.

Analog Multiplexer - The analog multiplexer (U26) is used by the CPU to monitor critical power supply voltages and the oxygen concentration (signal generated by the interface board). The CPU selects one channel at a time using the multiplexer channel select input.

Fans - There are three fan connections on the system board. These include the internal lithium ion battery cooling fan, motor cooling fan and exhaust fan mounted behind the power supply. The battery cooling fan and exhaust fan are controlled by the CPU while the motor cooling fan is controlled by the DSP.

Real Time Clock (RTC) - The RTC is set and read by the CPU using the I2C interface. Back-up power to the RTC is provided by a 20mm coin cell battery (B1) on the system board.

Cold Cathode Fluorescent Lamp (CCFL) - The power supply and brightness control circuit for the CCFL backlight inverter are on the system board (J20). The 12VDC supply is used to power the CCFL inverter and the CPU generates the PWM signal used to control the brightness of the CCFL backlight inverter using opamplifier (U20).

EEPROM - The system board contains a 32Kbit SPI controlled EEPROM (U8) for the storage of calibration and therapy data.

System LED - A blinking green led (CR14) is used to indicate normal system operation.

Vcc Supervisor - The microprocessor supervisory (U19) monitors the 3.3VDC DSP power supply. If the power supply falls below 3.08VDC the microprocessor supervisory will reset the DSP.

LCD - The CPU generates all the signals to control the LCD but the physical interface for the LCD is on the system board (J3).

JTAG - The DSP can be programmed via the JTAG interface (J2) on the system board or on the CPU board.



4.4 CPU DAUGHTER CARD

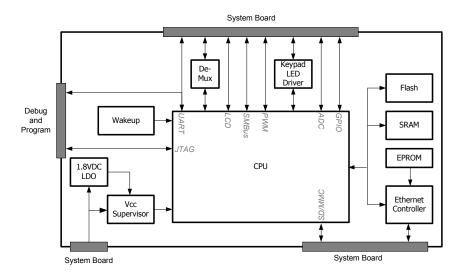


FIGURE 4-4 CPU DAUGHTER CARD BLOCK DIAGRAM

3.3VDC Supply - The CPU daughter card (except the core) operates on a 3.3VDC supply. This power is provided by the system board via the interface connector (J1).

1.8V Supply - The CPU has a core power requirement of 1.8VDC (250mA) generated by the fixed low-dropout linear voltage regulator (U5).

VCC Supervisor - A dual voltage supervisor (U10) monitors both the 3.3VDC and 1.8 VDC supplies. The thresholds for the supplies are set at 2.93V and 1.68V respectively. These thresholds preserve the system within the recommended operating ranges of the CPU, SDRAM and FLASH.

Central Processing Unit (CPU) - The Trilogy ventilator uses a 32-bit System-on-chip RISC Core CPU (U1). The processor is the brains behind all ventilator functions. Some of the key futures of the processor include the LCD controller, Synchronous and Asynchronous memory controller, MMC/SD controller, SMBUS interface and two UARTs.

There are two clock inputs used by the processor - a 14.7456 MHz crystal for system clock functions and a 32.768 kHz crystal for the RTC and power down modes. Even if the RTC is never used, the 32.768 kHz crystal must be connected to provide proper state transition.

Three external pins can generate a system reset. If any of these three lines is pulled low, a system reset will occur.

- 1. Power on Reset asserted by the dual VCC supervisor.
- 2. Power fail signal generated using pull-up resistor, and
- 3. User reset line connected to the system board through the slimstack connector.

The CPU provides the interface to the SD/MMC card. The physical interface for the SD/MMC card is a 6 wire interface (one signal, one clock and four data) on the system board.

Two of the three available UART ports of the Processor are used, one for the RS-232 communication through the system board connector while the other for debug operations through the debug connector.

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Wakeup - The wakeup circuit is a result of a unique requirement for the CPU. Approximately 1 to 2 seconds after the supply voltage reach a stable operating condition the CPU requires a rising edge on the wakeup signal to boot. To generate this wakeup signal a 555 timer (U9) is used. When power is applied the timer generates a 100Hz square wave output used as the wakeup signal for the CPU.

SDRAM - The CPU daughter card has a 64MB synchronous dynamic random access memory (SDRAM) (U2). This memory is controlled by the processor using the built-in SDRAM controller. This memory is used by the CPU as a "scratch pad". The SDRAM has a 13-bit address line (2-bit bank select and 11-bit address input line) and 32-bit data bus operates at 100MHz. The SDRAM is powered by the 3.3VDC supply.

Flash Memory The CPU daughter card has a 32MB nonvolatile, electrically block erasable (Flash) programmable memory (U3) configured for 16 bit word size. It stores the application code for the CPU. The flash memory is organized as thirty-two 128KB erase blocks and each block can be individually written and erased. The flash memory has a 22-bit address and 16-bit data bus. It is powered by 3.3VDC supply.

Ethernet Controller - The CPU daughter card provides a MAC to PHY interface using the Ethernet (non-PCI) controller (U6). This device is a mixed signal (Analog/Digital) device, that through the use of a 16 bit parallel interface generates the 4 necessary outputs signals for the 10/100 Mbps IEEE 8023 Ethernet communication. The PHY interface is a 4B5B/Manchester encoder/decoder. The Ethernet chip is powered by the 3.3VDC supply and uses an external clock input of 25MHz.

EERPOM - A serial EEPROM (U13) is connected to a SPI peripheral of the Ethernet controller. It is used by the controller to store default communication information and parameters.

Keypad LED Driver - The CPU uses a keypad led driver (U7) to provide eight additional I/O ports. The driver also provides higher output current capacity and to act as a possible ESD buffer for front panel LEDs.

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4.5 SENSOR BOARD

The sensor board is the main source of pressure and flow feedback for the Trilogy ventilator. It supplies the system board with real time digital pressure and flow data. This data is processed by the system board and used to control the blower speed and supply the prescribed pressure and flow support to the patient.

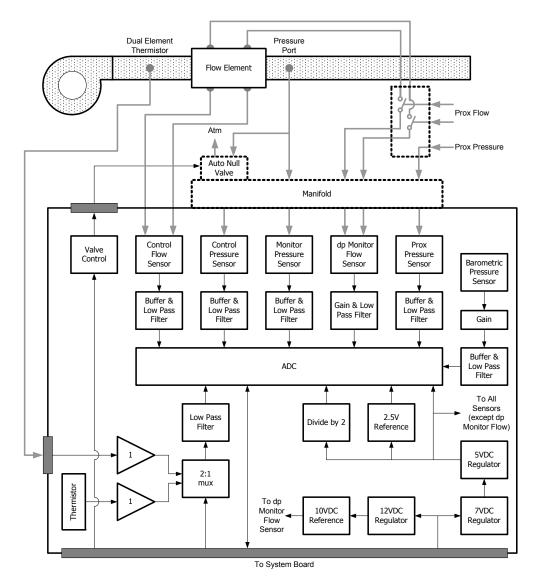


FIGURE 4-5 SENSOR BOARD BLOCK DIAGRAM

The circuitry on the board contains seven sensors, sensor conditioning circuits, ADC (Analog to Digital Converter), and linear and reference voltage sources. Six of the sensors are on board and one (the dual element temperature sensor) is located in the air stream and connected to the board through cabling (J1). Signals that are critical for patient therapy have separate sensors for control functions (machine blower control) and monitoring functions (alarms and patient parameters). All sensors are powered by the 5VDC regulator (U7) except dp monitor flow sensor (MT4) which is powered by the 10VDC precision reference (U10). All sensor outputs are ratio-metric to the power supply.

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Power Supply - The system board supplies the sensor board with bulk 14VDC (J4). The bulk 14VDC is regulated to 7VDC and 12VDC by voltage regulators U8 and U9 respectively. Precision reference (U10) generates the 10VDC using 12VDC to power the monitor flow sensor. Linear regulator (U7) generates the 5VDC using 7VDC to power the remaining sensor board with the exception of the solenoid valve.

The precision reference voltage regulator (U5) generates a 2.5VDC output signal used by the ADC (U10) and the instrumentation amplifier (U6). The ADCs internal reference gain amplifier generates a ratio-metric 4.096VDC reference voltage using this 2.5VDC voltage. The divide-by-two circuit generates a 2.5VDC signal. This signal is ratio-metric voltage of 5VDC supply. It is buffered and used as an input to the ADC (U2).

Pressure Sensors - The monitor pressure signal is measured using the monitor pressure sensor (MT1). It generates 0.2 to 4.7 VDC output for pressures range of 0 to 102 cm H20 pressure. The output of the sensor is buffered and filtered using an anti-aliasing real pole filter with a cutoff frequency set at 222 Hz. The proximal and control pressure signals are measured using the proximal and control pressure sensor (MT2 and MT3 respectively). They generate 0.5 to 4 VDC output for pressure range of 0 to 105 cm H20. The output of both the sensors is buffered and filtered using an anti-aliasing real pole filter with a cutoff frequency set at 222 Hz.

Flow Sensors - The monitor flow signal is measured using the monitor flow sensor (MT4). It generates a -25mv to +25mV output for pressures in the range of -10.5 to +10.5 cm H20. At the output of the sensor is amplified and filtered using the instrumentation amplifier (U6). The filter used is a 2 stage passive low pass filter with a cutoff frequency of 222Hz. The control flow signal is measured using the control flow sensor (MT5). It generates a 0.25 to 4VDC output for differential pressures in the range of -5 to +5 cm H20. The output of the device is a square root relationship with regard to voltage and dP. The output of the sensor is buffered and filtered using an anti-aliasing real pole filter with a cutoff frequency set at 222 Hz.

Barometric Pressure Sensor - The atmospheric pressure is measured using the barometric pressure sensor (MT6). The output of the sensor is scaled by a factor of .797 using a resistive divider network (R18 & R19). The output is then buffered and filtered using an anti-aliasing real pole filter with a cutoff frequency set at 222 Hz.

Temperature Sensor - A dual element temperature sensor measures the air stream temperature. This sensor is a negative temperature coefficient with nominal resistance of 10K @25°C. One output of the dual element temperature sensor is buffered and fed to the 2:1 multiplexer (U4). The other input is from the on board thermistor RT1 which has a negative temperature coefficient and a nominal resistance of 100k @ 25°C. The output of the multiplexer is filtered by a single pole low pass passive RC circuit. Based on the input to the 2:1 multiplexer from the system board, output of one thermistor is supplied to the ADC.

Analog to Digital Converter - The analog outputs of all these sensors are converted to digital signals using the 16 bit SAR (successive approximation register) ADC (U2). The ADC communicates with the system board via the 4 wire SPI bus interface (J4).

Plastic Manifold – The plastic manifold provides a mechanical pneumatic connection to the sensors and allows for mounting of the solenoid valve.

Solenoid Valve - The sensor board uses a solenoid valve to auto null/vent the control pressure sensor to atmosphere periodically via port 1 of the plastic manifold. The solenoid valve is externally mounted on the manifold and powered by a 29VDC nominal supply provided by the system board (J2). The N-Channel Power MOSFET (Q2) turns the solenoid valve ON-OFF.

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4.6 FRONT PANEL BOARD

The front panel board provides the user interface support in the Trilogy ventilator. It includes circuitry for the keypad, keypad backlight, audible alarm, visual alarm, and AC present indicator.

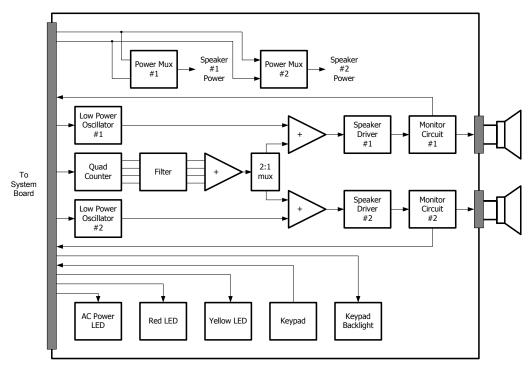


FIGURE 4-6 FRONT PANEL INTERFACE BOARD BLOCK DIAGRAM.

AC Power LED – A green LED (CR3) indicates the presence of AC/DC supply to the Trilogy ventilator.

Red LED – Two red LED's (CR2 & CR19) are used to indicate urgent and high priority alarm conditions. They are driven by two separate control lines from the CPU and the CPLD for redundancy during power fail conditions. Compared to normal operation, during power fail the red LED's have lower luminous intensity achieved using current limiting resistors (R105, R106) to preserve power consumption.

Yellow LED – Two yellow LED's (CR4 & CR5) are used to indicate low priority alarm conditions.

Keypad / User Input Switch Circuit – The front panel has 6 four-legged metal dimple types dome switches for user input. They support alarm silence, start/stop, left, right, up, and down functions. All share active low signal configuration. They are powered by the 3.3V power supply with the exception of the start/stop and right function keys which use the 3.3V_PLD to support additional function during the power fail and low power mode.

Keypad Backlight - An array of 12 white LED's (CR7 to CR18) are used to provide back light for the keypad. All the while LED's are controlled by a common switch (Q2).

Audible Alarm Signal - Each speaker has a mono-tone and a multi-tone audio alarm signal source. The mono tone audio alarm signal is generated by the low power oscillators (U3 & U5) running at a fixed frequency of 2.9 KHz. This frequency is set using external resistors (R12 & 14). The multi-tone audio alarm signal is generated using a combination of four harmonic signals. The 4-bit binary counter (U6) receives a square wave signal at various frequencies to generate these four harmonic signals. Each of the four harmonics signals is wave shaped using high-pass (RC) filters and low-pass (active) filter (U1). A summing amplifier (U2) then adds the

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four harmonic signals and generates the multi-tone audio alarm signal. Based on the input to the 2:1 multiplexer (U4) the multi-tone signal is fed to one speaker driver circuit at a time.

Audible Alarm Driver - The two audio alarm signals are fed to the respective piezo speaker drivers (U7 & U11) via the summing amplifiers (U14 & U15). The piezo speaker drivers have integrated boost converters and audio power amplifiers capable of driving the speaker load with a 16Vp-p signal. They provide pop-and-click removal function through bypass capacitors (C17 & C49) which determine the turn-on time of the audio amplifier. In addition, the speaker driver circuit provides the band switch function for a soft alarm sounding, and low power shutdown to support low power mode.

Speaker Monitoring Circuit - The operation of each speaker is verified using the current monitors sub-circuit. When a speaker is turned ON the high-side current shunt monitor (U8 & U12) generates an output current proportional to the differential voltage generated across the current sense resistors (R27 & R59). The output signal is used by the comparators (U9 & U17) to generate the alarm status signals for the CPU to verify speaker operation. The reference voltage for the comparators (U9 & U17) is set at 193mV.

Speaker Circuit Power - There are two independent power sources available to power the two speaker circuitry. The power mux (U13 & U16) select one of the two available power sources for each speaker circuitry. Both the speaker circuits use the different power supplies as primary and backup supplies.

4.7 INTERFACE BOARD

The interface board provides the means for the Trilogy ventilator to communicate with external devices such as the humidifier and other accessory devices. It includes circuitry for ethernet communication, RS-232 and RS-485 communication, oxygen sensing, remote alarm interface and accessory power generation.

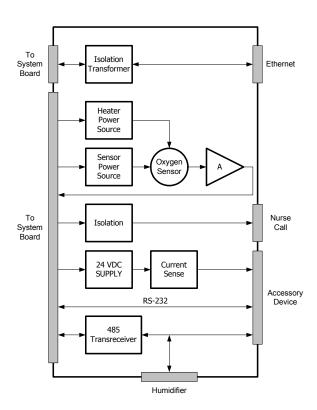


FIGURE 4-7 INTERFACE BOARD BLOCK DIAGRAM

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Ethernet - The Trilogy ventilator supports CAT-5 twisted pair based 10Mbs Ethernet communication protocol.

Oxygen Sensor - The Trilogy ventilator uses a ceramic zerconia solid-electrolyte oxygen sensor (U8) mounted on the interface board to detect hazardous level of oxygen leak inside the unit. A buck-boost switching regulator (U7) supplies a constant 1.5W power to the ceramic heater for oxygen sensor heater. A linear voltage regulator (U16) generates the 3.3 VDC supply for the oxygen sensor output gain amplifier (U10). In addition, it is also used to generate the 1.8VDC supply (U9) to power the oxygen sensing element. The power to the heater element and to the sensing element on the oxygen sensor can be disabled by using the switch (Q5). This helps to prolong the sensor life when not used. There is a minimum recommended delay of 3 minutes for the oxygen sensor to warm up and generate valid oxygen concentration measurements.

Remote Alarm (Nurse Call) Interface - The remote alarm sub-circuit generates the alarm signal for the Respironics remote alarm device and nurse call system. Electric isolation between the Trilogy ventilator and the remote alarm device is achieved using optical isolators (U3 & U6).

Regulated 24VDC supply - The interface board receives 29VDC bulk power supply from the system board and generates a regulated 24VDC supply using the linear voltage regulator (U1). The supply is used as power source for the accessory module and can be disabled by the system board using the switch (Q4). The high side current shunt monitor (U2) uses a current sense resistor (R14) to monitor the output current of the regulated 24VDC supply. If output current of the supply increases above 400mA the linear voltage regulator (U1) is shut down.

RS-232 Communication - The Trilogy ventilator supports standard full duplex RS-232 serial communication protocol. It is used for debug mode communications only.

RS-485 Communication - The Trilogy ventilator supports standard multi-drop half duplex RS-485 serial communication protocol. It is used by the Trilogy ventilator to communicate with the humidifier and accessory devices.

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CHAPTER 5: SYSTEM SETUP

NOTE

Please refer to the appropriate Provider and/or User Manual for additional information.

5.0 CHAPTER OVERVIEW

This chapter provides information regarding setup and operation of the Trilogy Ventilator as needed for servicing, repairing, and testing of the device.

5.1 KEYPAD LOCK FEATURE

There is a Keypad Lock feature that users can enable from the Options menu. It is intended to prevent accidental changes to device settings. This feature will lock the navigation keys (Up, Down, Left, and Right keys). If the Keypad Lock is enabled, a Keypad Unlock message will display on the bottom of the screen any time you press one of the navigation keys, as shown below.

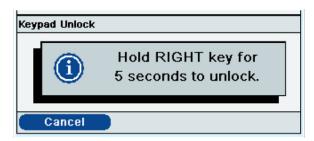


FIGURE 5-1: KEYPAD UNLOCK MESSAGE

If the keypad is locked, you must unlock it before you can enter the Menu. To unlock the keypad and enter the menu, you must first hold the Right button for 5 seconds to unlock the keypad. An audible indicator sounds when the keypad is successfully unlocked. Once the display is unlocked, you can enter the Menu as you normally would by pressing the Up button.

NOTE

- There is a keypad lock inactivity time-out period. After you have unlocked the keypad as indicated, the keypad will re-lock after five minutes of inactivity to prevent someone from accidentally pressing a button and changing any of the settings.
- When Keypad Lock is enabled, the Left, Right, and Up/Down buttons are locked while the ventilator is turned on. The Alarm Indicator/Audio Pause and Start/Stop buttons continue to function normally.
- The Keypad will automatically unlock if an alarm or informational message occurs and will remain unlocked the entire time alarms are active.
- Pressing the Left (Cancel) button will cancel the Keypad Unlock action.

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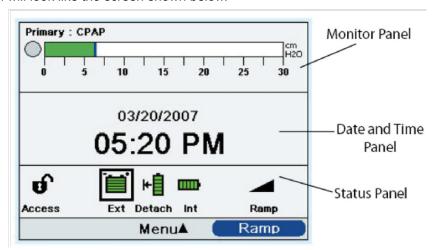
5.2 Accessing the Startup and Monitor Screens

1. After you press the On/Off button to begin therapy, the Startup screen below appears momentarily, indicating the device name and the software version.

NOTE

No Soft Buttons display on the Monitor screen if Keypad Lock is enabled.

2. The next screen that appears is the Monitor screen. The appearance of this screen will vary, depending on how you set up the device. If Detailed view is turned off in the options menu, your screen will look like the screen shown below.



NOTE

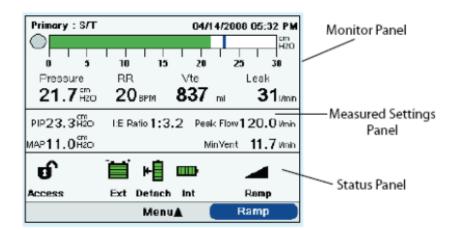
The symbols that appear on the Monitor Screen are described in detail later in this chapter.

- a. The top section of the screen, called the Monitor panel, shows the therapy mode and, if you set up a dual prescription for the patient, the Prescription indicator appears, indicating Primary or Secondary prescription. The patient breath symbol also displays during a patient-triggered breath, and a bar graph displays the current pressure level.
- b. The center section of the screen displays the current date and time.
- c. The bottom section, called the Status panel, displays certain symbols that indicate features being used, such as Ramp, as well as battery status.

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If Detailed view is turned on in the Options Menu, the Monitor screen will look like the screen shown below.



This screen contains more detailed information about the therapy.

- a. The top Monitor panel contains the Prescription indicator if a dual prescription exists, the therapy mode, a graph displaying the current pressure, and the current date and time. Additionally, this panel also displays patient pressure, respiratory rate (RR), exhaled tidal volume (Vte), and leak.
- b. The second panel in Detailed view is the Measured Settings panel. It provides patient-related data including Peak Inspiratory Pressure (PIP), Minute Ventilation, Peak Inspiratory Flow, Mean Airway Pressure (MAP), and I:E Ratio.
- c. The third panel is the Status panel and shows the same information displayed in the Detailed View Off screen, including features in use such as Ramp and battery status.

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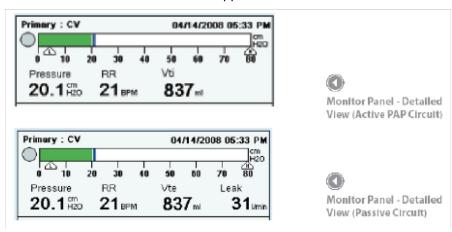
5.3 MONITOR SCREEN INDICATORS

This section describes the following indicators:

- Monitor Panel Indicators
- Measured Settings Panel Indicators
- Status Panel Indicators

5.3.1 MONITOR PANEL INDICATORS

All of the indicators that may appear on the Monitor Panel are described in detail in the following table. The Detailed view is shown below. The monitor panels for the Active circuit and Passive circuit are different, so both panels are shown. Some of these items do not appear in the Detailed View Off screen.



INDICATOR	DESCRIPTION
Prescription	If you set up a dual prescription for the patient, the words "Primary" or "Secondary" appear in the top left corner of the panel to indicate which prescription is active.
Therapy Mode	The current therapy mode displays at the top of the panel (for example, CPAP, S, S/T, etc.). If a special feature such as Flex, AVAPS, or Sigh is active, this feature will appear next to the therapy mode.
Date and Time	If you are in Detailed view, the current date and time appear in the top right hand corner of the panel. (In Detailed View Off, they appear in the center panel.)
Patient Breath	This symbol displays during a patient-triggered breath.

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INDICATOR	DESCRIPTION
Airway Pressure Manometer and Peak Pressure Symbol	The manometer (bar graph) displays the airway pressure in the patient circuit at all times. The manometer bar moves to the right as pressure increases during inhalation, and moves to the left as pressure decreases during exhalation. The peak pressure is also indicated on this bar. It is positioned according to the maximum patient pressure reached during each breath. The Peak Pressure symbol appears as a blue bar on the manometer. If a High Inspiratory Pressure alarm occurs, the Peak Pressure symbol changes from blue to red.
Low Pressure Indicator	If you enable a volume therapy mode, this indicator appears below the manometer bar, indicating the low pressure alarm setting.
High Pressure Indicator	If you enable a volume therapy mode, this indicator appears below the manometer bar, indicating the high pressure alarm setting.
Pressure	This indicator displays the current patient pressure. This only appears in detailed view.
Respiratory Rate (RR)	This indicator displays the measured respiratory rate in Breaths Per Minute (BPM). This only appears in detailed view.
Exhaled Tidal Volume (Vte)	This indicator displays the estimated exhaled tidal volume in milliliters and reflects compensation for BTPS. This only appears in detailed view when Passive Circuit is selected.
Inhaled Tidal Volume (Vti)	This indicator displays delivered tidal volume in milliliters and reflects compensation for BTPS. This only appears in detailed view when Passive Circuit is selected.
Leak	This indicator displays the total leak (non-returned flow), between the unit outlet and the patient, averaged over the previous breath. This only appears in detailed view when the Passive Circuit is selected.

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5.3.2 TRILOGY 100 & TRILOGY 200 MEASURED SETTINGS PANEL

All of the indicators that may appear on the Measured Settings panel (available only in Detailed view), are described in the following table.

INDICATOR	DESCRIPTION
PIP	Peak Inspiratory Pressure displays the maximum pressure delivered to the patient during the previous breath.
I:E Ratio	Displays a comparison of the time spent in inspiration to the time spent in expiration during the previous breath.
Peak Flow	Displays the maximum inspiratory flow delivered to the patient during the previous breath in I/min BTPS.
MAP	Displays the Mean Airway Pressure, which is the weighted average of pressure in the patient's airway over 6 breaths.
MinVent	Minute ventilation displays the amount of air delivered to the patient over the last minute in I/min BTPS.

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TRILOGY O2 & TRILOGY 202 MEASURED SETTINGS PANEL

All of the indicators that may appear on the Measured Settings panel (available only in Detailed view), are described in the following table.

INDICATOR	DESCRIPTION
PIP	Peak Inspiratory Pressure displays the maximum pressure delivered to the patient during the previous breath.
I:E Ratio	Displays a comparison of the time spent in inspiration to the time spent in expiration during the previous breath.
Peak Flow	Displays the maximum inspiratory flow delivered to the patient during the previous breath, reflecting compensation for BTPS.
MAP	Displays the Mean Airway Pressure, which is the weighted average of pressure in the patient's airway over 6 breaths.
FiO ₂	Displays the FiO ₂ setpoint.
MinVent	Minute ventilation displays the amount of air delivered to the patient over the last minute, reflecting compensation for BTPS.

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5.3.3 STATUS PANEL INDICATORS

All of the indicators that may appear on the Status Panel are described in the following table.



INDICATOR	DESCRIPTION
Ð	Indicates that the device is in Full Menu Access mode, which means you can adjust all prescription settings. Respironics recommends that you change the device to Limited Menu Access mode before giving the device to the patient, so patients cannot adjust their prescription settings. Only trained health care professionals and home care providers should adjust prescription settings.
	Displays when a Secure Digital (SD) memory card is inserted in the ventilator.
	Displays when the ventilator detects an error with the SD card.
	Displays at all times when an external battery is attached to the ventilator. The level of green shading shown in the symbol indicates the battery capacity and will go down as the battery charge level decreases. When the entire symbol is green, the battery is fully charged.
+	Displays at all times when a detachable battery is attached to the ventilator. The level of green shading shown in the symbol indicates the battery capacity and will go down as the battery charge level decreases. When the entire symbol is green, the battery is fully charged.
	Displays at all times, indicating the status of the internal battery. The level of green shading shown in the symbol indicates the battery capacity and will go down as the battery charge level decreases. When the entire symbol is green, the battery is fully charged.
	A black box displays around the battery that is currently supplying power to the ventilator when AC power is not available. (In the status panel shown above, the external battery is in use, so the symbol displays.)

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INDICATOR	DESCRIPTION
8	A yellow lightning bolt symbol displays with the Detachable or Internal battery symbol to indicate when the battery is charging. (In the status panel shown here, the detachable battery is being charged, so the symbol appears.
	Displays when the Audio Pause button has been pressed and Audio Pause is active. The alarm is silenced for one minute when the Audio Pause button is pressed.
	Displays when the Ramp feature is active.

NOTE

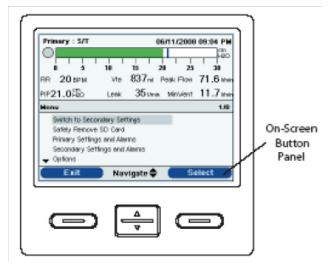
If a battery in use is very low (less than 20 minutes remaining), the inside of the box surrounding the battery symbol will change to yellow and all of the indicators in the battery will be empty. If a battery in use is near depletion (less than 10 minutes remaining), the inside of the box surrounding the battery symbol will change to red, and all of the bar indicators in the battery will be empty. These color changes only occur for the last available battery source.

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5.3.4 On-Screen Button Panel

The illustration below shows the on-screen button panel on the Main Menu screen, in relation to the buttons on the front of the Trilogy Ventilator.



NOTE

The on-screen buttons will vary depending on what screen you are in and what settings are enabled on your device.

At the very bottom of the display screen is the on-screen button panel. This panel corresponds with the control buttons on the ventilator.

- The left on-screen button specifies the action for the Left button on the device.
- The center on-screen button specifies the action for the Up/Down buttons on the device.
- The right on-screen button specifies the action for the Right button on the device.

5.3.5 NAVIGATING THE MENU SCREENS

To navigate through all of the menu screens and settings:

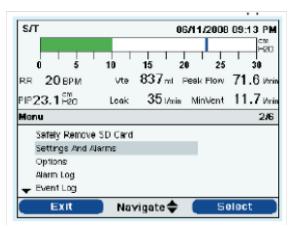
- Use the Up/Down button to scroll through the menu options and settings.
- Use the Left and Right buttons to perform the actions specified on the Left and Right on-screen buttons.

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5.4 CHANGING AND VIEWING SETTINGS IN FULL MENU ACCESS MODE

You can view and change settings using the Menu screens when the device is in Full Menu Access mode. To enter the Menu screens from the Monitor screen, press the Up button on the ventilator. The Main Menu screen shown below appears.



NOTE

- In the example Main Menu screen shown, the 2/6 that appears in the Menu banner indicates that item 2 is highlighted from a total of 6 items in the menu.
- For some therapy settings, once you reach the highest or lowest setting available, pressing the Up/Down button again will cycle back through the settings. For the parameters that do not wrap, when you reach the highest or lowest setting possible, a "Limit Reached" message appears in the Menu Banner on-screen.
- If you change a setting but decide you do not want to save it, you can press the Left button to cancel the change

Choose from the following selections on the Main Menu screen:

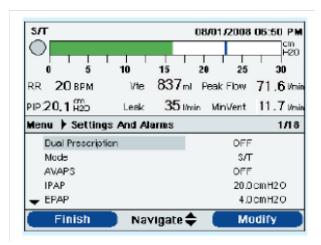
- Safely Remove SD Card: This option will appear if an SD card is inserted in the ventilator. Select
 this option when you want to remove the SD card. When the "Remove SD Card" confirmation
 message appears, remove the card. If you press the left (cancel) button or don't remove the card
 within 60 seconds, the confirmation message will close and the ventilator will continue writing to
 the card.
- Settings and Alarms: View and change prescription settings and alarms.
- **Options**: View and change device settings, such as Full or Limited Access mode, Detailed View, Language, etc.
- Alarm Log: View a list of the 20 most recent alarms that have occurred.
- **Event Log:** View a list of all events that have occurred, such as ventilator setting changes, ventilator inoperative conditions, alarms, etc.
- **Information:** View detailed information about the device, such as the device's software version and serial number.

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5.5 CHANGING THE DEVICE SETTINGS AND ALARMS

From the Main Menu screen, use the Up/Down button to highlight the Settings and Alarms menu, and press the Right button to select the menu. A screen similar to the one below appears.



5.5.1 DEVICE SETTINGS COMMON TO ALL THERAPY MODES

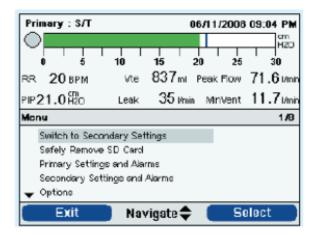
Some of the settings on this menu will vary depending on the therapy mode you select. The section below describes all of the settings that are common to all therapy modes.

DUAL PRESCRIPTION SETTING

You can turn the dual prescription setting on or off. Enable the setting if you want to create two separate prescriptions for the patient. For instance, you may want to set up a daytime prescription and then a separate nighttime prescription. If you enable this setting, then the menu options on the Main Menu will change to include new options:

- Switch to Primary/Secondary Settings
- Primary Settings and Alarms
- Secondary Settings and Alarms

The Main Menu screen will look like the screen below.



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5.5.2 MODE SETTING

You can change the Mode setting on the Setting and Alarms screen to one of the following therapy modes:

- CPAP Continuous Positive Airway Pressure
- S Spontaneous Ventilation
- S/T Spontaneous/Timed Ventilation
- T Timed Ventilation
- PC Pressure Control Ventilation
- PC-SIMV Pressure Controlled Synchronized Intermittent Mandatory Ventilation
- CV Control Ventilation
- AC Assist Control Ventilation
- SIMV Synchronized Intermittent Mandatory Ventilation

5.5.3 CIRCUIT TYPE

There are two types of circuit types you can select:

- Passive
- Active PAP
- Active Flow (Trilogy 200, Trilogy O₂, & Trilogy 202 Only)

The Passive circuit uses the Whisper Swivel II passive exhalation device. The Active PAP circuit type uses an active exhalation device with a proximal air pressure sensing connection. The Active Flow circuit type uses an active exhalation device with a proximal flow sensor.

When using the Passive circuit, the ventilator displays estimated patient pressures based on the resistance of the standard patient circuit (Whisper Swivel II with 1.8 meter tubing). Adding accessories to the patient circuit (humidifier, water trap, etc.) may cause an increase in circuit resistance and cause the device to display slightly higher pressures than what is actually delivered to the patient.

With the Active PAP or Active Flow circuit type selected, patient pressure is measured directly and is not affected by any change in circuit resistance.

The Passive circuit provides leak compensation. When using the Passive circuit in Volume Ventilation, the set Vti is delivered to the patient above the calculated circuit and cuff (or mask) leak. This is different from traditional active circuit ventilation where the cuff (or mask) leak reduces the tidal volume delivered to the patient. Volume ventilation with the Passive circuit delivers an inspiratory tidal volume close to the device setting regardless of leak; this should be considered when transitioning a patient from an active to a passive circuit. With a Passive circuit, Vte is estimated based on the calculated sum of circuit and cuff (or mask) leak.

The Active Flow circuit monitors proximal flow and proximal pressure. When using the Active Flow circuit configuration, flow trigger with leak compensation may be enabled. The default setting when using the Active Flow Circuit is Leak Compensation **On**. The clinician has the option to turn off the leak compensation;



however, unintentional leak will not be compensated. Both options measure the flow at the proximal flow sensor. Leak compensation is not available with PAP circuit configuration.

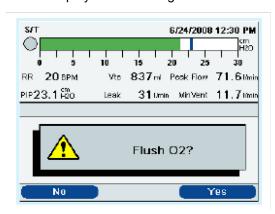
NOTE

- To change the circuit type, you must be in the Setup screen with the airflow turned off.
- When the Circuit Type setting is set to Passive Circuit, all Ramp Start Pressure settings in all modes will maintain the minimum range.
- When the Circuit Type setting is set to Active PAP Circuit, the Flex feature is unavailable.

5.5.4 FIO₂

You can set the Fractional Inspired Oxygen, which blends in a certain amount of oxygen into the air stream delivered to the patient. The amount of oxygen blended is based on this setting. You can set the FiO_2 from 21% to 100% in increments of 1.

A Flush feature is also available for the FiO_2 setting. A Flush button appears on-screen if the FiO_2 setting is greater than 21%. Pressing this button displays the following confirmation screen:



Selecting **Yes** allows you to temporarily increase the delivered oxygen concentration to 100% for two minutes. After two minutes is completed, the oxygen concentration will return to the previous FiO_2 setting. Selecting **No** cancels the action and does not changed the FiO_2 setting.

5.5.5 CIRCUIT DISCONNECT

WARNING

You should not rely on any single alarm to detect a circuit disconnect condition. The Low Tidal Volume, Low Minute ventilation, Low Respiratory rate, and Apnea alarms should be used in conjunction with the Circuit Disconnect alarm.

This setting enables or disables the circuit disconnect alarm. If enabled, an audible alarm will sound when a large, continuous air leak (such as mask removal) has been detected in the circuit.

You can choose **Off** to disable the alarm. Or, you can increase or decrease the setting from 10 to 60 seconds in 5 second increments. For example, a setting of 10 means that the alarm will sound after the circuit has been disconnected for 10 seconds.

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5.5.6 APNEA

This setting enables or disables the apnea alarm. If enabled, an audible alarm will sound when an apnea is detected.

You can choose **Off** to disable the alarm. Or, you can increase or decrease the setting from 10 to 60 seconds in 5 second increments. For example, a setting of 10 means that the alarm will sound if the time between spontaneous breaths exceeds 10 seconds.

5.5.7 Low VTE

This setting enables or disables the Low Vte alarm. The alarm activates when the estimated exhaled tidal volume is less than or equal to this setting. You can choose **Off** to disable this alarm, or you can increase or decrease the setting from 40 ml to 2000 ml in 5 ml increments. It cannot be set higher than the High Vte setting.

When AVAPS is **On**, the alarm activates when the calculated tidal volume is less that 90% of the target tidal volume setting. This alarm can be set to on or off.

5.5.8 HIGH VTE

This setting enables or disables the High Vte alarm. The alarm activates when the estimated exhaled tidal volume is greater than or equal to this setting. You can choose **Off** to disable this alarm, or you can increase or decrease the setting from 50 ml to 2000 ml in 5 ml increments. It cannot be set lower than the Low Vte setting, except to be turned off.

5.5.9 Low VTI

This setting enables or disables the Low Vti alarm. The alarm activates when the measured inhaled tidal volume is less than or equal to this setting. You can choose **Off** to disable this alarm, or you can increase or decrease the setting from 40 ml to 2000 ml in 5 ml increments. It cannot be set higher than the Vti setting.

5.5.10 HIGH VTI

This setting enables or disables the High Vti alarm. The alarm activates when the measured inhaled tidal volume is greater than or equal to this setting. You can choose **Off** to disable this alarm, or you can increase or decrease the setting from 40 ml to 2000 ml in 5 ml increments. It cannot be set lower than the Vti setting except to be turned off.

5.5.11 LOW MINUTE VENTILATION

This setting enables or disables the Low Minute Ventilation alarm. The alarm activates when the calculated minute ventilation is less than or equal to this setting. You can choose Off to disable this alarm, or you can increase or decrease the setting from 1l/min to 99 l/min in 1 l/min increments. It cannot be set higher than the High Minuted Ventilation setting.

5.5.12 HIGH MINUTE VENTILATION

This setting enables or disables the High Minute Ventilation alarm. The alarm activates when the calculated minute ventilation reaches or exceeds this setting. You can choose Off to disable this alarm, or you can increase or decrease the setting from 1l/min to 99 l/min in 1 l/min increments. It cannot be set lower than the Low Minute Ventilation setting except to be turned off.

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5.5.13 LOW RESPIRATORY RATE

This setting enables or disables the Low Respiratory Rate alarm. The alarm activates when the measured respiratory rate is less than or equal to this setting. You can choose Off to disable this alarm, or you can increase or decrease the setting from 4 BPM to 80 BPM in 1 BPM increments. It cannot be set higher than the High Respiratory Rate setting.

5.5.14 HIGH RESPIRATORY RATE

This setting enables or disables the High Respiratory Rate alarm. The alarm activates when the measured respiratory rate reaches or exceeds this setting. You can choose Off to disable this alarm, or you can increase or decrease the setting from 4 BPM to 80 BPM in 1 BPM increments. It cannot be set lower than the Low Respiratory Rate except to be turned off.

5.5.15 Low Inspiratory Pressure

This setting configures the Low inspiratory Pressure alarm. It is only user-settable in CV, AC, and SIMV modes. It cannot be set lower than PEEP + 2 pressure units or higher than the High Inspiratory Pressure. For passive circuits, you can increase or decrease the Low Inspiratory Pressure from 6 to 40 pressure units in increments of 1. For active circuits, you can increase or decrease the setting from 2 to 40 pressure units in increments of 1. For pressure modes, this alarm is not user-settable.

5.5.16 HIGH INSPIRATORY PRESSURE

This setting enables or disables the High inspiratory Pressure alarm. It is only user-settable in CV, AC, and SIMV modes. The High Inspiratory Pressure cannot be set lower than the Low Inspiratory Pressure. You can increase or decrease the High Inspiratory Pressure from 10 to 80 pressure units in increments of 1. For pressure modes, this alarm is not user-settable.

5.6 ADDITIONAL SETTING SPECIFIC TO THERAPY MODES

The Settings and Alarms menu also contains many additional settings specific to the various therapy modes. The specific settings for each therapy mode are described below.

5.6.1 CONTINUOUS POSITIVE AIRWAY PRESSURE (CPAP) MODE

In addition to the general settings described in the previous section of this manual, you can also set the following setting in CPAP mode.

1. CPAP

You can increase or decrease the CPAP pressure setting from 4 to 20 pressure units in increments of 1

NOTE

If the CPAP pressure is set to 4 (the minimum setting), the Ramp Length setting will be unavailable.

Trigger Type

The Trilogy can be set to trigger breaths based on automatic flow thresholds or specific flow settings. You can select either **Auto-Trak** or **Flow Trigger** as the Trigger Type. When Auto-Trak is selected, the AutoTrak trigger initiates based on automatic flow thresholds. When Trigger Type is



set to Flow Trigger, Flow Trigger Sensitivity and Flow Cycle Sensitivity become active, and the trigger initiates based on the Flow Trigger Sensitivity setting.

NOTE

Auto-Trak is only available if the Passive Circuit is selected.

Trigger Type is not available when Active PAP or Active Flow circuit is selected. Flow trigger is the triggering method used for Active PAP and Active Flow circuits.

3. Flow Trigger Sensitivity

If you set the Trigger Type to Flow Trigger, the Flow Trigger Sensitivity setting displays. You can increase or decrease the setting from 1 to 9 l/min in 1 l/min increments.

The flow trigger initiates when the patient's inspiratory effort creates a flow equal to or greater than the flow sensitivity setting

NOTE

Flow Trigger with leak compensation is only available if the Active Flow circuit is selected.

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4. Leak Compensation (Trilogy 200, Trilogy O2, & Trilogy 202 Only)

If you are using an Active Flow circuit, you can turn Leak Compensation On or Off.

NOTE

Enabling Leak Compensation when using the Active Flow Circuit configuration only affects triggering and does not affect tidal volume delivery or Vte measurement.

5. Flow Cycle Sensitivity

If you set the Trigger Type to Flow Trigger, the Flow Cycle Sensitivity setting displays. You can increase or decrease the setting from 10 to 40 percent (%) in 1% increments for Trilogy 100 and from 10 to 90 (%) in 1% increments for Trilogy 200, Trilogy O_2 , & Trilogy 202.

6. Ramp Length

The Ramp Length allows you to set the ramp time. You can disable Ramp by selecting Off, or you can increase or decrease the Ramp Length setting from 5 to 45 minutes in 5-minute increments.

NOTE

The Ramp Start Pressure setting will not display if the Ramp Length is set to Off or if the CPAP pressure is set to 4 pressure units.

7. Ramp Start Pressure

You can increase or decrease the ramp start pressure in increments of 1 to 4 pressure units to the CPAP pressure setting. The patient also has access to this setting, unless the ramp length is set to Off.

NOTE

Ramp Start Pressure is less than or equal to CPAP -1 pressure unit in CPAP mode.

8. Flex

You can enable or disable the Flex setting. Off disables the setting and prevents the patient from using Flex. To enable the setting, set Flex to 1, 2, or 3. The patient also has access to this setting, if Flex is enabled. The Flex feature is not available when using an active circuit.

NOTE

- In CPAP mode, Flex is only available when CPAP is greater than 4 pressure units.
- In S mode, Flex is only available when EPAP is greater than or equal to 4 pressure units and IPAP is less than or equal to 25 pressure units.



5.6.2 SPONTANEOUS (S) MODE

In addition to the general settings described in the previous section of this manual, you can also set the following setting in S mode.

1. AVAPS

AVAPS is only available if Flex is not enabled.

You can disable AVAPS by selecting **Off**, or you can enable AVAPS by selecting **On**. If you select Off, the IPAP setting displays. If you select On, the IPAP Max Pressure and IPAP Min Pressure display. These settings are described below.

NOTE

AVAPS is only available if the Passive Circuit is selected.

IPAP

NOTE

IPAP, IPAP Max, or IPAP Min cannot be set to more than 30 pressure units above EPAP.

The IPAP setting displays if AVAPS is Off. You can increase or decrease the Inspiratory Positive Airway Pressure (IPAP) from 4 to 50 pressure units in increments of 1. IPAP is limited to a maximum of 25 pressure units when Flex is enabled. You cannot set the IPAP setting lower than the EPAP setting.

3. IPAP Max Pressure

The IPAP Max Pressure setting displays if AVAPS is enabled. You can increase or decrease the setting from 4 to 50 pressure units in increments of 1. The IPAP Max Pressure must be equal to or greater than the IPAP Min value.

4. IPAP Min Pressure

The IPAP Min Pressure setting displays if AVAPS is enabled. You can increase or decrease the setting from 4 to 50 pressure units in increments of 1. The IPAP Min Pressure must be equal to or greater than the EPAP value, and it must be less than or equal to the IPAP Max Pressure.

5. EPAP

NOTE

EPAP cannot be set to more than 30 pressure units below IPAP, IPAP Max, or IPAP Min.

You can increase or decrease the Expiratory Positive Airway Pressure (EPAP) from 4 to 25 pressure units in increments of 1. For active circuits, EPAP can be set to zero.

When AVAPS is disabled, the EPAP setting must be less than or equal to the IPAP setting. When AVAPS is enabled, the EPAP pressure must be less than or equal to the IPAP Min Pressure.



6. Tidal Volume

The Tidal Volume setting displays if AVAPS is enabled. You can increase or decrease the setting from 50 to 2000 ml in 5 ml increments. Use this setting to establish the target volume of gas which the ventilator will produce and deliver during each Spontaneous breath.

NOTE

In CV, AC, and SIMV modes, the tidal volume setting is limited by the Inspiratory Time, to maintain the system's minimum and maximum peak flows.

7. Rise Time

NOTE

The Rise Time setting only displays if Flex is disabled. If the Flex is enabled, the device will use a rise time of 3.

You can adjust the rise time to find the most comfortable setting for the patient. Increase or decrease the setting from 1 to 6 until you find the right setting. The rise time of 1 to 6 corresponds to tenths of a second (e.g., a setting of 4 equals 0.4 second rise time).

8. Apnea Rate

NOTE

In S/T, T, PC, PC-SIMV, SIMV, CV, and AC modes, the Apnea Rate is greater than or equal to the Breath Rate and is limited by the current Inspiratory Time setting to maintain a minimum 1:1 I:E ratio.

If the Apnea alarm is enabled, you can set the Apnea Rate from 4 to 60 BPM in 1 BPM increments. In S mode, the Apnea Rate is greater than or equal to 1:2 I:E ratio.

5.6.3 SPONTANEOUS/TIMED (S/T) MODE

All of the settings described in the S Mode section are also available in S/T mode, except the Flex setting. In addition to those settings, the settings below are also available in S/T mode.

1. Breath Rate

NOTE

In volume modes, the Breath Rate range is limited by the current Inspiratory Time setting to maintain a minimum 1:1 I:E ratio.

In AC mode, you can increase or decrease the Breath Rate setting from 0 to 60 BPM, while in all other modes, you can increase or decrease the setting from 1 to 60 BPM in 1 BPM increments. Use the Breath Rate setting to establish the minimum rate of mandatory breaths that the ventilator will deliver per minute.

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2. Inspiratory Time

You can adjust the Inspiratory Time setting from 0.3 to 5.0 seconds in 0.1 second increments. Inspiratory Time is the duration for the inspiratory phase of a mandatory breath.

NOTE

- In pressure modes, the inspiratory time range is limited by the current Breath Rate setting to maintain a minimum 1:1 I:E ratio.
- In volume modes, the inspiratory time range is limited by the current Tidal Volume and Breath Rate settings to maintain a minimum 1:1 I:E ratio and the system's minimum and maximum peak flow.

5.6.4 TIMED (T) MODE

All of the settings available in S/T mode are available in T mode, except for the Trigger Type setting. Please refer to the descriptions in the S and S/T Mode sections of this chapter for detailed information on the T mode settings.

5.6.5 PRESSURE CONTROL MODE

All of the settings available in S/T mode are available in PC mode, except for the Flow Cycle Sensitivity setting (when Flow Trigger is enabled). Please refer to the descriptions in the S and S/T Mode sections of this chapter for detailed information on the PC mode settings.

5.6.6 Pressure Control Synchronized Intermittent Mandatory Ventilation (PC-SIMV) Mode

The following settings, described in the S and S/T mode sections of this section, also are available in PC-SIMV mode:

- Breath Rate
- Inspiratory Time
- Trigger Type
- Flow Trigger Sensitivity (if Trigger Type is set to Flow Trigger)
- Flow Cycle Sensitivity (if Trigger Type is set to Flow Trigger)
- Rise Time

In addition to these, the following settings are also available in PC-SIMV mode.

1. Pressure

You can increase or decrease the Pressure setting from 4 to 50 pressure units in increments of 1. This is the pressure the ventilator will deliver during the inspiratory phase of a mandatory or assist breath.



2. Pressure Support

You can increase or decrease the Pressure Support setting from 0 to 30 pressure units in increments of 1. This is the pressure support the ventilator will deliver during the inspiratory phase of a Spontaneous breath.

NOTE

In PC-SIMV mode, you cannot set up Pressure Support for Mandatory and Assist breaths (Pressure - PEEP) greater than 30 pressure units.

3. PEEP

The Positive End Expiratory Pressure (PEEP) setting can be increased from 0 to 25 pressure units in active circuits and 4 to 25 pressure units in passive circuits, in increments of 1. PEEP is the positive pressure maintained in the patient circuit during exhalation. The PEEP must be less than or equal to the pressure setting.

NOTE

The Pressure Support and PEEP settings together cannot exceed 50 pressure units.

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5.6.7 CONTROL VENTILATION (CV) MODE

The following settings, described in the previous sections of this chapter, are also available in CV mode:

- Tidal Volume
- Breath Rate
- Inspiratory Time
- PEEP
- High Inspiratory Pressure
- Low Inspiratory Pressure

NOTE

The Low Inspiratory Pressure is limited to PEEP +2 in CV, AC, and SIMV modes.

In addition to these, the following settings are also available in CV mode.

1. Flow Pattern

NOTE

The Flow Pattern setting might be limited to only Ramp or Square based on the Tidal Volume, Inspiratory Time, and Breath Rate settings to maintain the minimum and maximum peak flows.

You can choose either Ramp or Square for the Flow Pattern setting.

2. Sigh

You can enable or disable the Sigh setting by selecting **On** or **Off**. A Sigh is a breath that is delivered every 100 breaths at 150% of the normal volume.

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5.6.8 ASSIST CONTROL (AC) MODE

The AC mode contains the following settings described in the S, S/T, PC-SIMV, and CV mode sections in this chapter. Please refer to the descriptions in those sections for detailed information.

- Tidal Volume
- Breath Rate
- Inspiratory Time
- Flow Pattern
- PEEP
- Trigger Type
- Flow Trigger Sensitivity
- Sigh
- · High Inspiratory Pressure
- Low Inspiratory Pressure

NOTE

Flow Cycle Sensitivity is not available in AC mode.

5.6.9 SYNCHRONIZED INTERMITTENT MANDATORY VENTILATION (SIMV) MODE

NOTE

The High Inspiratory Pressure is greater than or equal to PEEP + Pressure Support in SIMV mode.

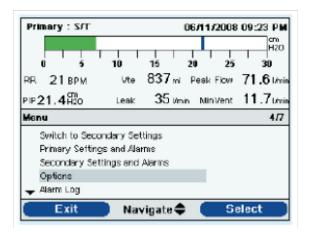
The SIMV mode contains the following settings described in the S, S/T, PC-SIMV, and CV mode sections in this chapter. Please refer to the descriptions in those sections for detailed information.

- Tidal Volume
- Breath Rate
- Inspiratory Time
- Pressure Support
- Flow Pattern
- PEEP
- Trigger Type
- Sigh
- Rise Time
- High Inspiratory Pressure
- Low Inspiratory Pressure

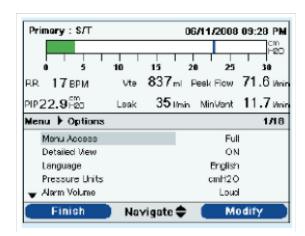


5.7 VIEWING AND CHANGING OPTIONS MENU ITEMS

From the Main Menu screen, select the Options item, as shown below.



The Options menu appears, shown in the screen below.



The following settings are available on the Options menu.

Menu Access – You can select Full or Limited menu access. Full menu access allows operators
to access all ventilator and prescription settings. Limited menu access allows operators to access
only certain settings and does not allow them to change prescription settings. To prevent patients
from tampering with prescription settings, do not give them Full menu access.



• **Detailed View –** You can turn Detailed View on or off using this setting. Detailed view displays additional settings and therapy information on the Monitor screen. The figures below show examples of the Monitor screen with Detailed View on or off.

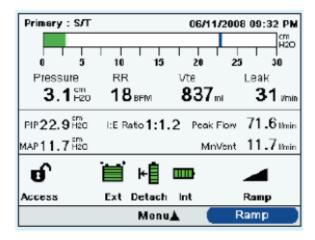


FIGURE 5-1: DETAILED VIEW ON

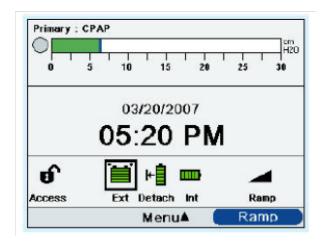


FIGURE 5-2: DETAILED VIEW OFF

- Language The next item on the Options menu allows you to select the Language that the software will appear in (English, French, German, etc.). The information on the screens will display in the language selected here.
- **Pressure Units –** The next item allows you to select the pressure units that will display on the screens. You can choose either:
 - cmH2O
 - hPa
 - mBar

All pressure units on the screens will display in the unit of measure selected here.

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Alarm Volume – You can adjust the volume of the Trilogy alarms using this setting. Select either
Loud or Soft as the alarm volume options.

WARNING

Make sure the alarm volume is sufficiently louder than the room noise level to ensure that the audible alarm can be heard by the caregiver.

- Keypad Lock You can enable or disable the Keypad Lock feature, which is described in detail
 earlier in this section. Enabling the Keypad Lock feature can prevent someone from accidentally
 pressing a button and changing any of the settings. Select On to enable the feature or Off to disable it.
- Keypad Backlight The next item you can set is the Keypad Backlight. You can turn the backlight
 On or Off using this setting. Whenever you press the On/Off button to begin therapy, the keypad
 backlight temporarily lights up. Once therapy is being provided, the keypad will be lit according to
 this Keypad Backlight setting. If the setting is On, the backlight remains on while therapy is provided. If the setting is Off, the backlight remains off while therapy is provided.
- LCD Brightness The LCD display is lit by a backlight. The backlight turns on when the initial Startup screen displays. You can adjust the brightness of the LCD backlight from 1 10, with 1 being the dimmest setting and 10 being the brightest.
- **Screen Saver –** You can change the screen saver to reduce power consumption or dim the screen in a dark room. You can choose the following settings:
 - Off: No screen saver displays and the LCD backlight remains lit at your brightness setting.
 - Breath: The display appears as a black screen, with only the patient breath indicator and manometer visible.
 - Black: The display's backlight is turned off and the display is black with no information visible.
 - Dim: The display's backlight is decreased, so that the display is still visible but not as bright.

If enabled, the screen saver will display after 5 minutes of no keypad activity. Pressing any button on the device will exit the screen saver. Also, any alarm or informational message will exit the screen saver.

NOTE

Setting the screen saver to Black allows the device to run for a longer period of time on battery power.

- **Date Format** You can choose either mm/dd/yyyy or dd/mm/yyyy as the date format that will display on the Trilogy screens.
- **Time Format** You can choose to display either an AM/PM time format or 24 Hour time format (for example, 2:49PM or 14:49).
- **Month –** The month defaults to the current month. The adjustable range is from 1 (January) 12 (December).
- **Day –** The day defaults to the current day. The adjustable range is from 1 31. The maximum value is based on the selected month.
- Year The year defaults to the current year. The adjustable range is from 2000 2099.



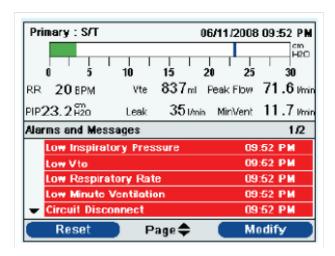
- Hour The hour defaults to the current hour. The adjustable range is from 12AM 12PM or 0-23, depending on the selected Time Format.
- Minute The minute defaults to the current minute. The adjustable range is from 0 59.
- **IP Address Mode** You can change the IP address mode to either DHCP or Static, depending on the type of network you are using (if applicable).
- Operational Hours The operational hours displays the total number of hours that the Trilogy blower has been on since the last time this value was reset. You can reset this value to zero (0) if desired (e.g., each time you give the device to a new patient). This value helps you determine how often the patient is using the device. The Operational Hours shown here differs from the Blower Hours shown on the Information screens. The Blower Hours displayed in the Information screen is the total number of hours that the blower has been working over the life of the device. You cannot reset this value.

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5.8 VIEWING THE ALARM LOG

From the Main Menu screen, you can select Alarm Log to access the Alarm Log screen. An example is shown below.



NOTE

In the Alarm Log screen, the 1/2 shown in the Menu banner indicates that page 1 of 2 alarm pages is being viewed at this time.

The alarm log displays the alarms in chronological order with the most recent events displayed first. It lists the 20 most recent alarms or messages that appeared on the Trilogy display. When the Trilogy is in Limited Menu access mode, the alarm log cannot be cleared. It can be cleared when in Full Menu access mode. Depending on how many alarms have occurred, the alarm log may be several pages long. The entries in the alarm log use the same names that displayed when the alarm initially occurred and was displayed in the Alarm View.

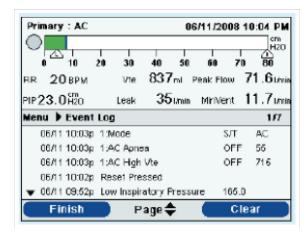
NOTE

In Full Menu access mode, you can press the Right (Clear) button to clear the alarm log if desired.



5.9 VIEWING THE EVENT LOG

From the Main Menu screen, you can select Event Log to access the Event Log Screen. An example is shown below.



The event log displays a list of all events that have occurred, such as ventilator setting changes, ventilator inoperative conditions, alarms, etc., in chronological order with the most recent events displayed first. When the Trilogy is in Limited Menu access mode, the event log is not available. It can be viewed and cleared when in Full Menu access mode. You can page through the event log if it is multiple pages. The number of pages appears in the upper right corner of the panel (in the example above, shown as 1/6).

In the event log descriptions, any description beginning with a 1: or 2: is a prescription change event. The 1 represents a change to a primary setting and the 2 represents a change to a secondary setting. This is followed by the setting that was changed.

The last two columns indicate setting and alarm changes. If the entry is a setting change, the first column shows the old setting value and the last column shows the new setting. If the entry is an alarm, the first column shows the value that triggered the alarm and the last column shows the number of seconds that the alarm was active.

NOTE

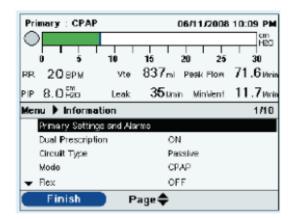
In Full Menu access mode, you can press the Right (clear) button to clear the event log if desired.

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5.10 VIEWING DEVICE INFORMATION

From the Main Menu screen, you can select Information to access the Information screen. You can also view the Information screen by holding the Down key for 5 seconds. This causes the detailed view of the Monitor Screen and the Information Menu to be displayed temporarily. This key sequence is valid from the Monitor Screen while in Full or Limited Access. An example is shown below.



The Information screen provides you with a summary of the current prescription settings, device settings, and system settings. You can use the Up/Down buttons to scroll through the information.



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CHAPTER 6: TROUBLESHOOTING & ALARMS

6.0 CHAPTER OVERVIEW

This chapter identifies the alarms associated with the Trilogy Ventilator. This chapter should be used to help service technicians diagnose problems with the Trilogy Ventilator, along with determining what parts, if any, need to be replaced.

6.1 TRILOGY VENTILATOR ALARMS

There are three types of alarms:

- High Priority Require immediate response by the operator.
- Medium Priority Require prompt response by the operator.
- Low Priority Require operator awareness. These alarms alert you to a change in the ventilator status.

Additionally, the ventilator also displays informational messages and confirmation alerts that notify you of conditions that need attention but do not qualify as alarm conditions.

NOTE

If multiple alarms occur at the same time, all alarms are processed and displayed, but the alarms are ordered first by priority and then by occurrence, with the newest, highest priority alarms at the top of the list. The alarm precedence is in the following order: high priority, medium priority, low priority, and informational messages.

NOTE

Not all alarms are available in every therapy mode; some alarms are mode dependant.

6.2 AUDIBLE AND VISUAL ALARM INDICATORS

When an alarm condition occurs:

- The alarm LED indicator on the Audio Pause button lights
- The audible alarm sounds
- A message appears on the screen describing the type of alarm
- · The remote alarm (if applicable) is activated

Each of these is described in detail below.

6.2.1 ALARM LED INDICATORS

The Alarm Indicator/Audio Pause button on the front of the ventilator lights up as follows whenever an alarm is detected:

- Red Flashing Indicator When the device detects a high priority alarm, the Alarm Indicator/Audio Pause button flashes red.
- Yellow Flashing Indicator When the device detects a medium priority alarm, the Alarm Indicator/ Audio Pause button flashes yellow.



• Yellow Solid Indicator - When the device detects a low priority alarm, a solid yellow light appears on the Alarm Indicator/Audio Pause button.

The Alarm Indicator/Audio Pause button does not light up when informational messages or confirmation alerts display.

6.2.2 AUDIBLE INDICATORS

NOTE

For the alarm indicators noted throughout this manual, each "dot" represents an audible beep.

An audible indicator sounds whenever a power failure or a high, medium, or low priority alarm is detected. Additionally, an audible indicator sounds for informational messages and to confirm that certain actions have occurred (for example, when an SD card is inserted or removed from the device).

- **Ventilator Inoperative Audible Indicator** When a ventilator inoperative alarm occurs, a continuous audible alarm sounds. The alarm descriptions later in this chapter display this indicator as:
- Power Failure Audible Indicator When a power failure occurs, a series of beeps sounds in a 1 beep pattern, repeating one second on, then one second off. The alarm descriptions later in this chapter display this indicator as:
- **High Priority Audible Indicator** When a high priority alarm is detected, a series of beeps sound in the following pattern, which is repeated twice: 3 beeps, a pause, and then 2 more beeps. This indicator continues until the cause of the alarm is corrected or the audible alarm is paused. The alarm descriptions later in this chapter display this indicator as: • • • • • •
- **Medium Priority Audible Indicator** When a medium priority alarm is detected, a series of beeps sound in a 3-beep pattern. This pattern repeats until the cause of the alarm is corrected or the audible alarm is paused. The alarm descriptions later in this chapter display this indicator as:
- Low Priority Audible Indicator When a low priority alarm is detected, a series of beeps sound in a 2-beep pattern. This pattern repeats until the cause of the alarm is corrected or the audible alarm is paused. The alarm descriptions later in this chapter display this indicator as:
- Informational Messages and Confirmation Audible Indicators When an informational message appears on screen, a brief, 1- beep audible indicator sounds. Additionally, when the device detects that a certain action has been completed (for example, when the Start/Stop button is pressed to start therapy, or when an SD card is inserted or removed from the device) a brief, 1-beep audible indicator sounds. The alarm descriptions later in this chapter display this indicator as: •

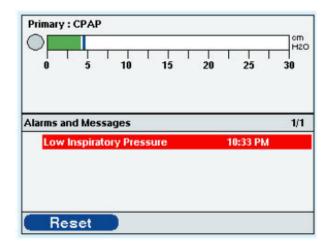


6.2.3 ALARM MESSAGES

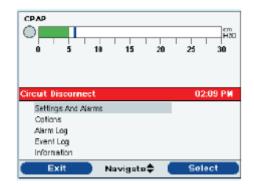
NOTE

An alarm message will also display in the Menu Banner if a menu is active when an alarm occurs.

When the ventilator detects an alarm, the Alarms and Messages Screen is displayed showing a description of the alarm condition. When an alarm message appears, it will be highlighted in red if it is a high priority alarm or in yellow if it is a medium or low priority alarm. (The highlight color matches the alarm LED color on the Alarm Indicator/Audio Pause button.) If an alarm is manually reset by the user, the Alarms and Messages screen is removed and the Monitoring Screen is re-displayed. If the alarm self-cancels, the Alarms and Messages screen remains displayed, but the highlight from the active alarm is removed, the LED is unlit, and the audible alarm stops. The screen below is an example of a possible alarm message.



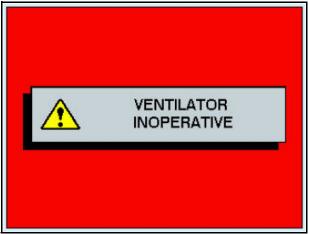
If a menu is displayed on the screen when an alarm occurs, the description of the newly generated alarm is displayed in the menu banner area. This is done so that the modification to the current parameter can be completed before addressing the alarm condition in case the modification affects the alarm condition. The screen below is an example of an alarm message displayed in the menu banner.



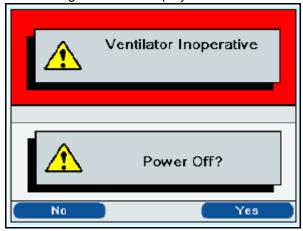
The Alarms and Messages Screen will automatically display in place of the Monitor screen when exiting from the menu system using the Exit soft key when an alarm is displayed in the menu banner. If an alarm is manually reset by the user or self-cancels, the menu banner on-screen before the alarm occurred will reappear.



If a Ventilator Inoperative alarm occurs, the entire display screen turns red and the Ventilator inoperative message appears on-screen, as shown below.



To turn the ventilator off from a Ventilator Inoperative condition, use the normal power off sequence. When the Start/Stop button is selected, the following screen will display.



Select the Right button (Yes) to turn the ventilator off and stop the audible alarm. Selecting the Left button (No) will return the screen to the Ventilator Inoperative Alarm Screen without silencing the audible alarm.

6.2.4 REMOTE ALARM

When the ventilator detects an alarm condition, if you are using a remote alarm system, a signal is sent from the ventilator to activate the remote alarm.

6.3 AUDIO PAUSE AND ALARM RESET FEATURES

This section describes the Audio Pause and Alarm Reset features.

6.3.1 AUDIO PAUSE

When an alarm occurs, you can temporarily silence the audible indicator by pressing the Alarm Indicator/Audio Pause button. The alarm is silenced for 60 seconds and then will sound again if the cause of the alarm has not been corrected. Each time the Alarm Indicator/Audio Pause button is pressed, the alarm silence period resets to one minute.

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When Audio Pause is active, the Alarm Indicator/Audio Pause symbol appears if you are on the Monitor screen. Additionally, an "Audio Pause" message displays in the menu banner on the Alarm Display screen.

You can 'pre-silence' alarms that have not yet occurred by pressing the Alarm Indicator/Audio Pause button while no alarms are active. Then, if an alarm occurs, the audible indicator does not sound until the Audio Pause time limit has expired.

6.3.2 ALARM RESET

The Reset button is used to clear the currently active alarm(s) from the display and stop the LED and audible alarm indicator. This button should be selected after the situation causing the alarm(s) has been corrected. All active alarms are cancelled and alarm detection is restarted when this button is selected.

The ventilator self-cancels certain alarms if the cause of the alarm is corrected, shutting off the alarm LED, the audible alarm, and the alarm background color. However, the alarm text remains on the screen. You can manually reset an alarm by pressing the Left button (Reset). The Audio Pause function is cancelled if the alarm is manually reset.

NOTE

- Alarm Reset will not affect ventilator inoperative alarms.
- When the device automatically resets an alarm, the alarm message remains displayed until you select the Left (Reset button).

6.3.3 ALARM VOLUME CONTROL

You can adjust the Alarm Volume from the Options menu. You can select Loud or Soft, depending on your preference.

WARNING

Make sure the alarm volume is set loud enough to be heard by the caregiver.

6.3.4 What to Do When an Alarm Occurs

WARNING

You should not rely on any single alarm to detect a circuit disconnect condition. The Low Tidal Volume, Low Minute Ventilation, Low Respiratory Rate, and Apnea alarms should be used in conjunction with the Circuit Disconnect alarm.

Complete the following steps when an alarm occurs:

- 1. Look at the alarm indicators and listen to the audible alarm sound. Note the color of the Alarm Indicator/Audio Pause button (red or yellow) and whether the LED is solid or flashing.
- 2. Look at the display to check the alarm message that appears on-screen and whether it is highlighted in red or yellow.
- 3. Press the Alarm Indicator/Audio Pause button to temporarily silence the audible alarm. A visual indicator displays if you are on the Monitor screen or an "Audio Pause" message appears in the menu banner on the Alarm Display screen.



4. Look up the alarm in the alarm descriptions later in this section to determine the source of the alarm and the appropriate action.

6.4 ALARM SUMMARY TABLE

The following table summarizes all of the high, medium, and low priority alarms and informational messages.

ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
Loss of Power	High	• •	It may occur when a complete power failure has occurred and power was lost while the device was providing therapy. This may happen if the internal battery was the only power source in use and was completely depleted.	Red flashing button; Blank screen	Shuts down
Ventilator Inoperative	High		It occurs when the ventilator detects an internal error or a condition that may affect therapy.	Red flashing button; "Ventilator Inoperative" message	Shuts down if can't provide therapy safely. Or, continues to operate at a limited level

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ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
Ventilator Service Required	High		It occurs when the device cannot perform to specification, a backup safety feature is compromised, or the delivery of therapy is compromised. If the problem is not corrected, the device will generate a reminder message once per hour, or whenever power is cycled, until the issue is corrected. Additionally, if the device is powered off, a reminder message will immediately appear when the device is turned on again.	Red flashing button; "Ventilator Service Required" message	Operates
Check Circuit	High	• • • • •	It occurs when the device detects a problem with the patient circuit, such as pinched or detached tubing, water condensation in the proximal pressure lines, or problems with the active exhalation device.	Red flashing button; "Check Circuit" message	Operates
Low Circuit Leak	High	• • • • •	It occurs when the system detects a problem with the leak device in the passive circuit.	Red flashing button; "Low Circuit Leak" message	Operates



ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
High Expiratory Pressure	High	•••	It occurs when the delivered pressure exceeds the target patient pressure during the expiratory phase by 5 pressure units. This may be due to pinched tubing or the patient having a fast breath rate. The alarm will automatically terminate when the delivered pressure comes within 5 pressure units during the expiratory phase.	Red flashing button; "High Expiratory Pressure" message	Operates
Low Expiratory Pressure	High	• • • •	It occurs when the delivered pressure is 5 pressure units or more below the target patient pressure during the expiratory phase. The alarm will automatically terminate when the delivered pressure comes within 5 pressure units of the target patient pressure during the expiratory phase.	Red flashing button; "Low Expiratory Pressure" message	Operates
High Internal Oxygen	High	• • • • •	It occurs when there is a leak in the internal air delivery system that allows oxygen to build up inside the device. The alarm is generated when the internal oxygen concentration reaches 5% above ambient levels.	Red flashing button; "High Internal Oxygen" message	Continues to operate when internal oxygen concentra tion reaches 5% above ambient levels.

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ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
High Oxygen Flow (Trilogy O ₂ & Trilogy 202 Only)	High	•••	It occurs when the concentration of oxygen from the device is 10% above the FiO ₂ set point for more than 30 seconds. This could be caused by a problem with the output of the oxygen source.	Red flashing button; "High Oxygen Flow" message	Operates
Low Oxygen Flow (Trilogy O ₂ & Trilogy 202 Only)	High	• • • •	It occurs when the concentration of oxygen from the device is 10% below the FiO ₂ set point for more than 30 seconds. This could be caused by the oxygen source being disconnected from the device, an occlusion in the tubing from the oxygen source to the device, or a problem with the output of the oxygen source.	Red flashing button; "Low Oxygen Flow" message	Operates
High Oxygen Inlet Pressure (Trilogy O ₂ & Trilogy 202 Only)	High	• • • • •	It occurs when the pressure of the oxygen from the source measures greater than 87 psi.	Red flashing button; "High Oxygen Inlet Pressure" message	Operates



ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
Low Oxygen Inlet Pressure (Trilogy O ₂ & Trilogy 202 Only)	High	•••	It occurs when the pressure of the oxygen from the oxygen source measures less than 40 psi. This could be caused by the oxygen source being disconnected from the device, an occlusion in the tubing from the oxygen source to the device, or a problem with the output of the oxygen source.	Red flashing button; "Low Oxygen Inlet Pressure" message	Operates
Circuit Disconnect	High	• • • •	It occurs when the breathing circuit is disconnected or has a large leak. The alarm will automatically terminate when the leak is fixed for 6 seconds and breaths are delivered again.	Red flashing button; "Circuit Disconnect" message	Operates
Apnea	High	• • • •	It occurs when the patient has not triggered a breath within the time specified in the apnea alarm setting. The alarm will automatically terminate when two consecutive patient breaths are detected that meet the apnea alarm time setting.	Red flashing button; "Apnea" message	Operates

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ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
High Vte	High	••••	It occurs when the estimated exhaled tidal volume is greater than the High Vte setting. The alarm will automatically terminate when a breath occurs in which the estimated exhaled tidal volume does not reach the High Vte alarm setting for three consecutive breaths.	Red flashing button; "High Vte" message	Operates
Low Vte	High	• • • • •	It occurs when the estimated exhaled tidal volume is less than the Low Vte alarm setting. The alarm will automatically terminate when a breath occurs in which the estimated exhaled tidal volume exceeds the Low Vte alarm setting for three consecutive breaths.	Red flashing button; "Low Vte" message	Operates
High Vti	High	• • • •	It occurs when the delivered tidal volume is greater than the High Vti alarm setting. The alarm will automatically terminate when a breath occurs in which the delivered tidal volume does not reach the High Vti alarm setting for three consecutive breaths.	Red flashing button; "High Vti" message	Operates



ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
Low Vti	High	• • • • •	It occurs when the delivered tidal volume is less than the Low Vti alarm setting. The alarm will automatically terminate when a breath occurs in which the delivered tidal volume exceeds the Low Vti alarm setting for three consecutive breaths.	Red flashing button; "Low Vti" message	Operates
High Respiratory Rate	High	• • • •	It occurs when the respiratory rate is greater than the High Inspiratory Rate alarm setting. The alarm will automatically terminate when the measured respiratory rate is less than the High Respiratory Rate alarm setting.	Red flashing button; "High Respiratory Rate" message	Operates
Low Respiratory Rate	High	• • • •	It occurs when the respiratory rate is less than the Low Respiratory Rate alarm setting. The alarm will automatically terminate when the measured respiratory rate is greater than the Low Respiratory Rate alarm setting.	Red flashing button; "Low Respiratory Rate" message	Operates

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ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
High Inspiratory Pressure	High	• (for first two consecutive occurrences) • • • (for 3rd consecutive occurrence) • • • • (for 10th consecutive occurrence)	It occurs in several stages and escalates from an audible beep for the first two occurrences to a high priority alarm if the problem continues. It is detected differently for volume and pressure therapy modes. For volume modes, the alarm will sound if the measured patient pressure exceeds the High Inspiratory Pressure setting specified by the clinician. The alarm will automatically terminate when the peak inspiratory pressure is less then or equal to the High Inspiratory Pressure alarm setting. For pressure modes, the alarm occurs when the delivered pressure exceeds the target patient pressure by 5 pressure units or more during the inspiratory phase. The device will automatically cycle to the expiratory phase and continue to operate. The alarm will automatically terminate when the delivered pressure falls within 5 pressure units of the target patient pressure during the inspiratory phase	Peak Pressure symbol turns red. When condition first occurs, a beep will sound. When condition occurs for third time, button flashes yellow and yellow "High Inspiratory Pressure" message appears. When condition occurs for 10th time, button flashes red and red "High Inspiratory Pressure" message appears.	Operates



ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
Low Inspiratory Pressure	High		It is detected differently for volume and pressure therapy modes. For volume modes, the alarm will sound if the measured patient pressure is less than the Low Inspiratory Pressure setting by the clinician. The alarm will automatically terminate when the peak pressure at the end of the breath is greater than or equal to the Low Inspiratory Pressure alarm setting. For pressure modes, the alarm occurs when the delivered pressure is 5 pressure units or more below the target patient pressure during the inspiratory phase. the alarm will automatically terminate when the delivered pressure comes within 5 pressure units of the target patient pressure during the expiratory phase.	Red flashing button: "Low Inspiratory Pressure" message	Operates
High Minute Ventilation	High		It occurs when the patient's minute ventilation is greater than the High Minute Ventilation alarm setting. The alarm will automatically terminate when the calculated minute ventilation is less than the High Minute Ventilation alarm setting.	Red flashing button; "High Minute Ventilation" message	Operates



ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
Low Minute Ventilation	High	•••	It occurs when the patient's minute ventilation is less than the Low Minute Ventilation alarm setting. The alarm will automatically terminate when calculated minute ventilation is greater than the Low Minute Ventilation alarm setting.	Red flashing button; "Low Minute Ventilation" message	Operates
Low Battery	Escalates from Medium to High	(Medium - when approx. 20 minutes remain) (High - when approx. 10 minutes remain)	It occurs when the last battery available is low or nearly depleted. This alarm occurs in two stages. When approximately 20 minutes of battery run time remains, a medium priority alarm is generated. If no action is taken and the battery continues to deplete, the alarm escalates to a high priority alarm when approximately 10 minutes of battery run time remains.	Medium Priority - Yellow flashing button. "Low Detachable Battery," "Low External Battery," or "Low Internal Battery" message appears in yellow, On Status Panel, box around battery is yellow High Priority - Red flashing button. "Low Detachable Battery," "Low External Battery," or "Low Internal Battery" message appears in red, On Status Panel, box around battery is red	Operates



ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
High Temperature	Escalates from Medium to High	• • • for medium • • • • • for high	It occurs when the estimated patient airstream temperature or the ventilator internal temperature is too high. The alarm occurs in several stages. Internal fans are started when the medium priority alarm is generated. If the condition causing the high temperature is not corrected and the temperature continues to rise, the alarm will escalate to the high priority alarm.	Yellow flashing button and yellow "High Temperature" message appears. If condition worsens, button flashes red and red "High Temperature" message appears.	Operates
Replace Detachable Battery	Low or High, depending on cause of alarm	• • for low • • • • • • • for high	It occurs when the detachable battery is nearing the end of its useful life or a failure in the detachable battery that prevents it from charging or discharging has been detected. The alarm occurs in several stages, from low to high priority. If the alarm is reset without removing the battery, the alarm will be regenerated once every hour until the detachable battery is removed. The detachable battery is not used, and the power source is switched to the next available power source if the alarm is the high priority alarm.	"Replace Detachable Battery" message appears. If battery is nearing end of useful life, message appears with yellow background and button is solid yellow. If battery fails, message appears with red background and button flashes red.	Operates



ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
Ventilator Service Recommended	Medium		It occurs when the device has detected an error, but the error will not affect device performance or safety. Therapy and safety are not compromised. If the problem is not corrected, the device will generate a reminder message once per day, or whenever power is cycled, until the issue is corrected. Additionally, if the device is powered off, a reminder message will immediately appear when the device is turned on again.	Yellow flashing button; "Ventilator Service Recommended" message	Operates
Keypad Stuck	Low	• •	It occurs when a key becomes lodged inside the case of the device.	Solid yellow button; "Keypad Stuck" message	Operates
Battery Discharging Stopped Due to Temperature	Info	•	This info message occurs when the detachable or internal battery becomes overheated while providing power for the device. The battery is not used and the power source is switched to the next available power source.	"Batt Discharge Stopped - Temp." message	Operates



ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
Battery Not Charging Due to Temperature	Info	•	This info message occurs when the detachable or internal battery becomes too hot while charging or the device was in too cold an environment before charging started. Battery charging stops until the battery cools or warms sufficiently.	"Batt Not Charging - Temp." message	Operates
Battery Not Charging	Info	•	This info message occurs when the device has detected an error condition with the battery that prevents it from accepting a charge. Battery charging stops.	"Detach Battery Not Charging" or "Internal Battery Not Charging" message	Operates
Check External Battery	Info	•	This info message occurs when a bad connection exists to the external battery or the external battery failed. The device continues to operate using power from the detachable battery, if available, or the internal battery.	"Check External Battery" message	Operates
Battery Depleted	Info	•	This info message occurs when the affected battery is fully depleted. The device continues to operate using the next available power source.	"External Battery Depleted" or "Detachable Battery Depleted" message appears	Operates



ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
AC Power Disconnected	Info	•	This info message occurs when the AC power source was lost, and the device has switched to an alternate power source (either a detachable or external battery, if connected, or the internal battery if no other source is available). If AC power returns, the ventilator will beep, but no message will appear on the display.	"AC Power Disconnected" message, and a box appears around battery in use.	Switches to alternate power source
External Battery Disconnected	Info	•	This info message occurs when the external battery power source is lost and the device has switched to an alternate power source (either a detachable battery, if connected, or the internal battery if no other source is available). If external power returns, the ventilator will beep, but no message will appear on the display.	"External Batt Disconnected" message, and a box appears around battery in use.	Switches to alternate power source



ALARM	PRIORITY	AUDIBLE	DESCRIPTION OF ALARM	VISUAL(ALARM INDICATOR BUTTON AND DISPLAY)	DEVICE ACTION
Detachable Battery Disconnected	Info	•	This info message occurs when the detachable battery power source is lost and the device has switched to an alternate power source (the internal battery if not other source is available). If detachable battery power returns, the ventilator will beep, but no message will appear on the display.	"Detachable Batt Disconnected" message, and a box appears around battery in use.	Switches to alternate power source
Start On Battery	Info	•	This info message indicates that the ventilator has started on battery power and no AC power is available. The device operator should verify that this is what is wanted.	"Start On Battery" message appears.	Operates
Card Error	Info	•	This info message occurs when an unusable SD card is inserted into the ventilator. The device continues to operate but data cannot be logged onto the SD card.	"Card Error" message appears	Operates

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6.5 TROUBLESHOOTING TABLE

DEVICE ACTION	POSSIBLE CAUSES	CORRECTIVE ACTION	
Unit does not power on	No power available	Verify AC or Batteries are present	
	 Front Panel Board On/Off circuit faulty 	Replace the Front Panel PCA	
	 System/CPU PCA does not recognize keypress 	Replace the System/CPU Subassembly PCA	
Unit power on, but Blower will not start	Blower not connected to PCA	Check connection from Blower to PCA	
	• Blower faulty	Replace Blower	
	Faulty motor drive on PCA	Replace System/CPU Subassembly PCA	
Unit is on, but Display is	 LCD not connected properly 	Check LCD Cable	
blank	 Inverter not connected properly 	Check Inverter and connection	
	Faulty Driver on System/CPU PCA	Replace System/CPU Subassembly PCA	
Unit and Display are on, but AC Power LED and	Front Panel not connected properly	Check connection from Front Panel to System.	
Keypad are not functioning	 Faulty Front Panel board 	Replace Front Panel Board	
Audible Alarm does not	Speakers not connected properly	Check Speaker connections	
beep 3 times at startup	 Faulty Speakers 	Replace Speakers	
	 Faulty Front Panel PCA 	Replace Front Panel PCA	
	Faulty System/CPU PCA	Replace System/CPU Subassembly PCA	
Any of the red (~1.5 sec), yellow (~1.5 sec), and	Front Panel not connected properly	Check connection from Front Panel to System	
white (~3 sec) LEDs do not light during startup	Faulty Front Panel board	Replace Front Panel Board	
Display flashes on and off	Bad power source	Verify good power source	
sporadically	CPU resetting	Replace System/CPU Subassembly	
Unit does not detect Detachable Battery	Internal connections not connected properly	Check connection on Power Management Board	
	Faulty Detachable Battery	Replace Detachable Battery	



Unit does not detect Internal Battery	Battery not connected properlyFaulty Internal Battery	Check connection on Power Management Board Replace Internal Battery
Unit does not detect External Lead Acid battery	 Battery not connected properly internally and/or externally Faulty External Lead Acid battery 	Check connections on Power Management Board and on outside of the unit Replace External Lead Acid battery
One alarm has much lower pitch or tone than other alarm	Faulty SpeakerFaulty Front Panel PCA	Replace Speaker Replace Front Panel Board
Display blank, but backlight working	Dislodged LCD (ribbon) cableFaulty LCD (ribbon) cable	Verify LCD cable/connections Replace LCD Cable
High Pos Flow at Calibration	Flow Straightener damaged or missing	Ensure Flow Straightener is installed in Flow Sensor Assembly correctly and is not damaged.
Trilogy Error Code relating to Battery issues	 Unit needs rebooted Erroneous read by Charger chip from power sequencing 	 Reboot unit using RASP command or allow unit to go to sleep Remove all power from unit, which includes physically disconnecting the Internal Battery, then reconnect.
Internal Battery fails current test or does not go into requested mode	 Faulty Detachable Battery Faulty Power Management Board 	 Replace Detachable Battery if found faulty. Replace Power Management Board.
Detachable Battery fails current test or does not go into requested mode	 Faulty External Battery Faulty Power Management Board 	Replace External Battery. Replace Power Management Board.
Unit won't reach Pressure Therapy settings in Run-in	 Incorrect settings Tubing setup incorrectly 	 Verify settings. Verify tubing is connected properly and there are no leaks. Verify Active Exhalation Valve is functioning properly.

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Failures involving pressure or flow tests	Tubing or porting block issuesFaulty Sensor PCA	 Verify tubing and porting block are connected properly and not leaking. If possible, verify faulty Sensor PCA, replace if suspected
Solid alarm when using Lead Acid or AC power fail alarm with known good Li- ion's	 CPU possibly rebooting or not running 	 Remove all power to reset and if possible obtain error logs; replace System/CPU PCA if suspected
Vent Service Required before software upgrade	• E143 (Pressure Span) in Error Log	Upgrade software. If error persists, or is not E143, refer to the proper Error Code Section of this chapter.
Trilogy error code relating to battery issues	Device software communication to the batteries	 Evaluate test report and/or Trilogy error log. If there is nothing conclusive from the evaluation, retest the unit.
High Pressure Leak Fail - this is indicative of a leak on the O2 flow side (Trilogy O ₂ & Trilogy 202 Only)	 Verify seating of the O₂ proportional valve Verify connection of green tube to manifold Verify green tube to port clip adaptor Verify O-rings on port clip adapter to pressure sensor are present Pressure sensor leaking 	 Reseat as needed or replace manifold if suspect. Reseat or replace green tube/ port clips as needed. Reseat or replace O-rings as needed. Replace PCA if all other related parts were verified not to leak.



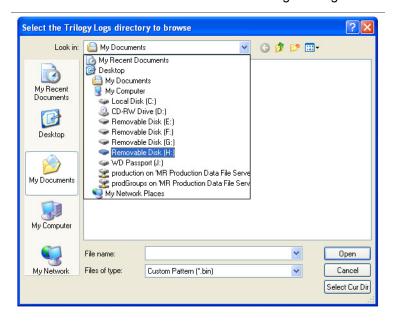
Low Pressure Leak Fail - This is indicative of a leak on the air flow side (Trilogy O ₂ & Trilogy 202 Only)	 Verify Interface between air duct and lower portion of air element seated properly Verify air element to mixer is intact and O-rings are not rolled Verify mixer to O₂ element is intact and O-rings are not rolled Verify O₂ element to manifold is intact and O-rings are not rolled Verify seating of the O₂ proportional valve Verify sense tubes from both O₂ and air elements to associated flow sensors are seated properly and free from holes or defects Ensure tubes are connected properly 	 Reseat or replace as needed. Replace manifold if suspect. Reseat all connections and retest. 	
Valve Functional Fail - Indicative of valve not properly opening or impeded air path (Trilogy O ₂ & Trilogy 202 Only)	 Verify seating on the O₂ proportional valve Foreign material in air path Faulty wire connection from O₂ proportional valve to board Faulty OBM PCA 	 Reseat as needed or replace manifold. Remove material if possible, otherwise replace manifold if suspect. Reseat connection or replace suspect part. Verify connector is seated, otherwise replace PCA 	
PCA Communication Fail - Indicative of PCA not communicating (Trilogy O ₂ & Trilogy 202 Only)	 Faulty OBM to Trilogy Serial Cable Faulty PCA 	 Verify cable connection, reseat or replace as needed Verify connector integrity. Reseat Cable at PCA Replace PCA. 	
Fan Test Fail - Indicative of a fan now working (Trilogy O ₂ & Trilogy 202 Only)	 Faulty OBM Fan Connection between fan and board Faulty PCA 	 Verify fan cable connection. Reseat connection. Replace Fan. Replace PCA 	
Current Test Fails - Indicative of idle current not within limits (Trilogy O ₂ & Trilogy 202 Only)	• Faulty PCA	Reseat connectionsReplace PCA	

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6.6 DOWNLOADING TRILOGY ALARM ERROR CODE INFORMATION

- 1. Insert a SD Card into the SD Card slot on the Trilogy Ventilator.
- 2. Press down the Audio Pause and Down Button at the same time.
- 3. Scroll down in the menu until you come to 'Write Event Log to SD Card". Select OK.
- 4. After write is complete, scroll back up to and select "Safely Remove SD Card"
- 5. Remove the SD Card from the Trilogy Ventilator.
- 6. Connect an SD Card Reader to your PC.
- 7. Place the SD Card into the SD Card reader.
- 8. Double click on the Trilogy Tool Box Icon.
- 9. Click on the BrowseUUTLogs Menu.
- 10. Select the location of the SD Card or PC Folder containing the Log Downloads.

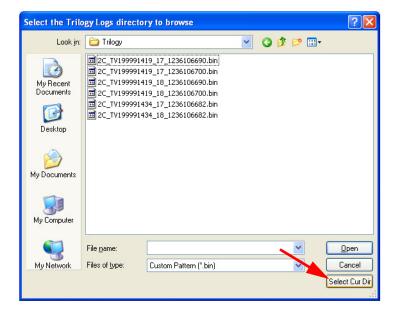




11. Select the Trilogy Folder.



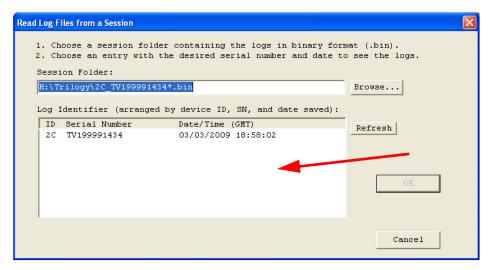
12. At the screen below, select the "Select Cur Dir" button. Do not select individual files.



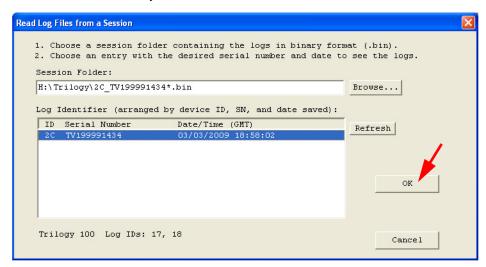
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13. The UUT for which you wish to view the files, will appear in the "Log Identifier" window.

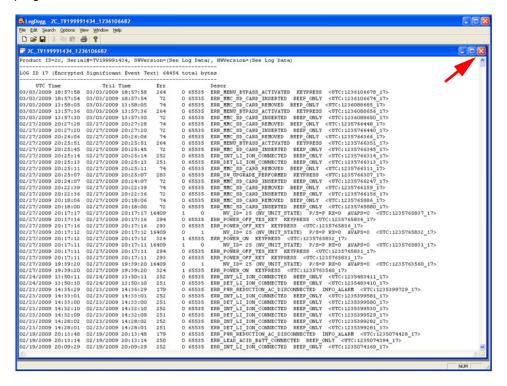


14. Select the file for the UUT you wish to view and then click on the "OK" button.





15. The encrypted files will then be displayed in the window. To close the file, select the red "X" in the top right corner of the window.



16. Select "No" to exit the program.



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6.7 TRILOGY ERROR CODES AND CORRECTIVE ACTIONS

NOTE

Use the corrective action(s) listed in section 6.5 (Troubleshooting Table) before using corrective action(s) listed in the error code charts below.

6.7.1 VENTILATOR INOPERATIVE ALARM ERROR CODES

ERROR CODE	PROBABLE CAUSE	CORRECTIVE ACTION
E-9	 Motor Capacitor not installed Motor Connector not soldered properly Motor Rotor Locked or other mechanical problem 	 Check the Capacitor Check Motor connector Replace Motor Replace System/CPU PCA
E-14	 Board revision resistors are not installed Software is running on wrong revision of the board 	Load Correct Software Replace System/CPU PCA
E-21	Problem with serial communication between DSP and Host	Replace System/CPU PCA
E-37	Host CPU is constantly rebooting	Reinstall Software Replace System/CPU PCA
E-71	Parameter Settings corrupted	Replace System/CPU PCA
E-73	Parameter Settings corrupted	Replace System/CPU PCA
E-76	Defective EEPROM	Replace System/CPU PCA
E-77	 Defective EEPROM Program Execution Error	Replace System/CPU PCA
E-90	 Software does not support newer/older revision of Hardware Board revision resistors on System/CPU PCA are setup incorrect 	Reinstall software Replace the System/CPU PCA
E-101	Program Execution Error	Contact Respironics Product Support
E-103	Control Flow Sensor out of electrical spec	Check TubingCheck ManifoldReplace Sensor PCA



E-125	3 reboots occurred within 24 hours	 Examine error logs for reasons for reboots Proceed accordingly
E-141	Motor connector not soldered correctly Motor Rotor Locked or other mechanical problem	 Check Bulk Capacitor Check Motor connector Replace Motor Replace System/CPU PCA
E-145	 Both control and Monitoring Pressure sensors failed. ADC failed 	Check tubingCheck manifoldReplace Sensor PCA
E-146	 Both control and Monitoring Pressure sensors failed. ADC failed 	 Check tubing Check manifold Replace Sensor PCA
E-158	• A serious alarm has occurred before six breaths have been taken after blower is turned on. The lack of therapy data causes back up therapy not to be initiated and the ventilator to become completely inoperative. The alarms that would initiate this possible scenario are: both pressure sensors failing, both pressure and flow sensors failing, a failure of non-volatile memory, or a high pressure patient alarm	 None - recorded for informational purposes to indicate that backup therapy could not be provided for the Ventilator Inoperative condition currently active
E-160	 Obstructed intake Obstructed flow path Obstructed or disconnected pressure tubing Faulty Blower 	 Check Inlet filter / Air Path for obstruction Check circuit Check internal tubing Replace Blower
E-163	Control Flow Sensor out of electrical specification	Check tubingCheck manifoldReplace Sensor PCA



E-172	 Unit is exceeding 60cm of delivered pressure High pressure sensor reading Large amount of drift Pinched/blocked tubing 	 Check the tubing for pinched or blocked tubes Check the circuit for leaks Check the Active Exhalation Valve functionality Check the tubing for leaks, kinks, or blockages Replace the Sensor board Replace the System/CPU PCA
E-253	 Communication failure The Host CPU is unable to communicate with the DSP 	Replace the System/CPU PCA
E-306	 Disconnected O₂ tubing O₂ level greater than 35% 	 Connect / replace O₂ tubing Replace Interface Board
E-323	When the device encounters errors while it has started to apply the Rx Setting from the SD Card, the device stops the Rx update and tries to restore back the old setting on the device. During the restoration of these old settings of the device, if there is an error this error code is generated	Replace System/CPU Subassembly
E-357	Control Flow Sensor out of electrical spec	Check tubingCheck manifoldReplace Sensor Board PCA

6.7.2 VENTILATOR SERVICE REQUIRED ALARM ERROR CODES

ERROR CODE	PROBABLE CAUSE	CORRECTIVE ACTION
E-17	Big Motor Capacitor not installed or wrong value or at the end of life cycle	 Install or Replace Capacitor Replace System/CPU PCA
E-31	 Problem with +3.3V_DSP U23 failure DSP failure 	Replace System/CPU PCA
E-32	 Faulty Sensor Board Cable Faulty Sensor Board Circuitry 	 Check Sensor Board Cable Replace Sensor Board PCA Replace System/CPU PCA



E-36	a Could not an an an and	a Devilege Overlage (ODL) DOA
E-30	Cpld not programmed Div 45 of 1/00 about all	Replace System/CPU PCA
	Pin 15 of U30 shorted	
	Cpld failed	
E-64	Unit not calibrated	Recalibrate the device
	Calibration Table corrupted	Replace the System/CPU PCA
E-65	Unit not calibrated	Recalibrate the device
	Calibration Table corrupted	Replace the System/CPU PCA
E-66	Unit not calibrated	Recalibrate the device
	Calibration Table corrupted	Replace the System/CPU PCA
E-67	The sensor has drifted so much that the positive or negative displacement of the zero point has moved the maximum or minimum calculable pressure value up or down somewhat. Enough positive displacement can reduce the top end of pressure readings below that expected by the Trilogy design. By similar reasoning, enough negative displacement can result in a condition where the pressure sensor cannot read low enough for Trilogy	Replace Sensor Board PCA
	The drift data for a sensor is also collected at zero pressure, and is converted to engineering units. When the absolute value (since the sensor can drift in either direction) of this drift exceeds a configured limit, a range error has occurred	
E-69	 The sensor has drifted so much that the positive or negative displacement of the zero point has moved the maximum or minimum calculable pressure value up or down somewhat. Enough positive displacement can reduce the top end of pressure readings below that expected by the Trilogy design. By similar reasoning, enough negative displacement can result in a condition where the pressure sensor cannot read low enough for Trilogy The drift data for a sensor is also collected at zero pressure, and is converted to engineering units. When the absolute value (since the sensor 	Replace Sensor Board PCA
	can drift in either direction) of this drift exceeds a configured limit, a range error has occurred	



E-70	The sensor has drifted so much that the positive or negative displacement of the zero point has moved the maximum or minimum calculable pressure value up or down somewhat. Enough positive displacement can reduce the top end of pressure readings below that expected by the Trilogy design. By similar reasoning, enough negative displacement can result in a condition where the pressure sensor cannot read low enough for Trilogy	Replace Sensor Board PCA
	The drift data for a sensor is also collected at zero pressure, and is converted to engineering units. When the absolute value (since the sensor can drift in either direction) of this drift exceeds a configured limit, a range error as occurred	
E-80	The sensor has drifted so much that the positive or negative displacement of the zero point has moved the maximum or minimum calculable pressure value up or down somewhat. Enough positive displacement can reduce the top end of pressure readings below that expected by the Trilogy design. By similar reasoning, enough negative displacement can result in a condition where the pressure sensor cannot read low enough for Trilogy	Replace Sensor Board PCA
	The drift data for a sensor is also collected at zero pressure, and is converted to engineering units. When the absolute value (since the sensor can drift in either direction) of this drift exceeds a configured limit, a range error has occurred	
E-94	Barometric pressure sensor is reading maximum or minimum counts for an extended period of time	Replace the Sensor Board PCA
E-100	Software error Bad table written by Production or Field Service Calibration	Recalibrate the device Replace the System/CPU PCA
E-106	Speaker 1 failed or not installed	 Install speaker 1 Check connection between front panel and system board Replace the Front Panel PCA Replace the System/CPU PCA



E-107	Speaker 2 failed or not installed	 Install speaker 2 Check connection between front
		panel and system board
		Replace the Front Panel PCA
		Replace the System/CPU PCA
E-130	The sensor has drifted so much that the positive	Check tubing
	or negative displacement of the zero point has moved the maximum or minimum calculable	Check manifold
	pressure value up or down somewhat. Enough positive displacement can reduce the top end of pressure readings below that expected by the Trilogy design. By similar reasoning, enough negative displacement can result in a condition where the pressure sensor cannot read low enough for Trilogy	Replace the Sensor Board PCA
	The drift data for a sensor is also collected at zero pressure, and is converted to engineering units. When the absolute value (since the sensor can drift in either direction) of this drift exceeds a configured limit, a range error has occurred	
E-131	Control pressure sensor is reading maximum or	Check tubing
	minimum counts for an extended period of time	Check manifold
		Check the Sensor Board Cable
		Replace the Sensor PCA
		Replace the System/CPU PCA
E-132	Monitor pressure sensor is reading maximum or	Check tubing
	minimum counts for an extended period of time	Check manifold
		Check the Sensor Board Cable
		Replace the Sensor PCA
		Replace the System/CPU PCA
E-133	Proximal pressure sensor is reading maximum or	Check tubing
	minimum counts for an extended period of time	Check manifold
		Check the Sensor Board Cable
		Replace the Sensor PCA
		Replace the System/CPU PCA

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E-134	Control flow sensor is reading maximum or minimum counts for an extended period of time	 Check tubing Check manifold Check the Sensor Board Cable Replace the Sensor PCA Replace the System/CPU PCA
E-135	Monitor flow sensor is reading maximum or minimum counts for an extended period of time	 Check tubing Check manifold Check the Sensor Board Cable Replace the Sensor PCA Replace the System/CPU PCA
E-136	Air stream Temperature 1 sensor is reading maximum or minimum counts for an extended period of time	 Check Airstream Temp Sensor connections Replace Sensor Replace Sensor Board Replace System/CPU PCA
E-137	dP Pressure Temperature sensor is reading maximum counts for an extended period of time	 Check dP Pressure Temp Sensor Replace Sensor Replace Sensor Board PCA Replace System/CPU PCA
E-138	Barometric Pressure sensor is reading a count outside of its valid range	Replace the Sensor PCA Replace the System/CPU PCA
E-139	Software Error	Recalibrate the device Replace the System/CPU PCA
E-143	Control or Monitoring Pressure sensor has drifted too far away from the other	 Turn off Blower and allow 30 seconds for drift processing to complete Replace the Sensor PCA



E-147	 The sensor has drifted so much that the positive or negative displacement of the zero point has moved the maximum or minimum calculable pressure value up or down somewhat. Enough positive displacement can reduce the top end of pressure readings below that expected by the Trilogy design. By similar reasoning, enough negative displacement can result in a condition where the pressure sensor cannot read low enough for Trilogy The drift data for a sensor is also collected at zero pressure, and is converted to engineering units. When the absolute value (since the sensor can drift in either direction) of this drift exceeds a configured limit, a range error has occurred 	Replace Sensor Board
E-161	 Internal Li-Ion has a TDA due to Discharge Overcurrent or Discharge Short Circuit PMB fault System Board fault Internal Li-Ion fault 	 Replace Power Management PCA Replace Internal Li-Ion battery Replace System/CPU PCA
E-164	Monitor Flow Sensor out of electrical spec	Check tubingCheck manifoldReplace Sensor Board
E-174	 Can't communicate with Charger Chip PMB fault Internal Li-Ion fault Detachable Li-Ion fault 	 Replace Power Management PCA Replace Internal Li-Ion battery Replace Detachable Li-Ion battery Replace System/CPU PCA
E-175	 Can't communicate with Internal Li-lon battery after five tries and charger is not trying to wake-up charge the battery Could take up to 210 seconds for alarm to be reported because the Charger is trying to wake-up charge the Internal Li-lon PMB fault Internal Li-lon fault 	Replace Power Management PCA Replace Internal Li-Ion battery
E-177	 Internal Li-lon battery not Authentic Fake battery PMB fault Internal Li-lon fault 	Replace Internal Li-Ion battery Replace Power Management PCA



		<u></u>
E-189	 Detachable or Internal Li-Ion usable and V_battLi 9 volts for greater than 10 sec indicates Li-Ion battery unable to power the System PMB fault 	Replace Power Management PCA
E-191	 AFE chip does not receive the appropriate frequency on the WDI pin from the Gas Gauge chip. Both of these chips are i the Li-Ion battery pack. Internal Li-Ion fault 	Replace Internal Li-Ion battery Replace Power Management PCA
E-193	Internal Li-Ion battery permanent failure	Replace Internal Li-Ion battery
E-195	 Internal Li-Ion Battery State of Health ≤ 50%. Full charge capacity (FCC) is less than 51% of the Design Capacity Internal Li-Ion FCC error Internal Li-Ion fault Internal Li-Ion end of life 	 Cycle the Internal Li-Ion (discharge battery so that capacity change is > 50%, then rest for > 2 hours. Charge to 100%, then rest for > 2 hours) Replace Internal Li-Ion battery
E-197	 Internal Li-Ion not present as indicated by the Charger Chip status PMB fault Internal Li-Ion battery pack fault 	 Replace Power Management PCA Replace Internal Li-Ion battery
E-203	Circuitry failure	Replace System/CPU PCA
E-205	Error indicates a shorted Q25, Q26 in the battery discharge path on the Power Management PCA. If either FET is shorted and AC is connected, then AC voltage will be connected to the Pb Acid battery terminals if one is connected and enabled. This may bring down the AC supply due to excessive current because the AC supply is trying to charge the PB Acid. If the AC supply folds back the unit should run properly	Replace the Power Management PCA
E-207	Q11 failure on Power Management PCA	Replace Power Management PCA
E-208	 Internal Li-Ion Cycle Count exceeded ≥ 475 cycles Internal Li-Ion end of life 	Replace Internal Li-Ion battery
E-265	Checks if Li-Ion battery is present by attempting to communicate when the Charger CHip safety signal indicates the battery is not present. Safety signal from the Li-Ion may be disconnected or Charger CHip may have a fault so it is not detecting a Li-Ion battery	Replace the Internal Battery Replace the Power Management PCA



E-267	 FCC > 150% of the Li-Ion design capacity Corrupted Gas Gauge 	 Cycle the Internal Li-Ion (discharge battery so that capacity change is > 50%, then rest for > 2 hours. Charge to 100%, then rest for > 2 hours) Replace Internal Li-Ion battery
E-269	 Error indicates an open VBULK FET. This MOSFET connects the output of the AC/DC supply to the input of the boost converter. Unit will not run on AC power. Will use battery power if available 	Replace Power Management PCA
E-270	 Error indicates an open Q11, which controls the discharge of the Li-Ion batteries post VbattLi. When the Lithium Ion power is selected, this FET is turned on. If Q11 fails open, there will be no Li- Ion battery power to the system 	Replace Power Management PCA
E-272	Charger chip failureFET failure	Replace Power Management PCA
E-273	Unit not calibrated	Recalibrate the device
	Calibration table corrupted	Replace the System/CPU PCA
E-274	Unit not calibrated	Recalibrate the device
	Calibration table corrupted	Replace the System/CPU PCA
E-290	One or more "Ventilator Service Required" errors are active in the system	 Examine the Significant Event Log error log. The first column of data contains the first error code that is causing this error message. Address according to this first error code
		Each time you address an error, cycle the motor (OFF to ON to OFF) and check the Significant Event Log again until this alarm is no longer sounded
		Remove all power from the unit and then reapply power, then turn the motor ON to see if this message is still reported
E-302	Voltage reference on internal ADC failed	Replace System/CPU PCA
E-311	 Unit wasn't calibrated or calibration table was destroyed 	Recalibrate Replace System/CPU PCA



E-333	U24 circuitry failureU26 circuitry failure	Replace System/CPU PCA
E-338	 Unit not calibrated Calibration Table corrupted	Recalibrate Replace System/CPU PCA
E-340	 CPLD stopped sending alive signal to the Host CPU 	Replace System/CPU PCA
E-344	Failure of Oxygen Blending Module	Recalibrate the device Replace the Blending Module PCA
E-349	Communication between Host and OBM was lost	Replace OBM PCA Replace Interface PCA
E-356	 Before a drift is computed for this sensor, its auto-null valve is opened to atmosphere. A test is performed to ensure that the valve has fully opened in accordance with the time provided by the technical specification 	Replace Sensor PCA Replace System/CPU PCA
E-358	Proximal Flow Sensor out of electrical spec.	 Check tubing Check manifold Replace Sensor PCA

6.7.3 VENTILATOR SERVICE RECOMMENDED ALARM ERROR CODES

ERROR CODE	PROBABLE CAUSE	CORRECTIVE ACTION
E-15	Hardware current limiting is not working correctly	Replace the System/CPU PCA Replace Motor
E-16	Problem with power management board or batteries (if no AC present)	Replace respective battery Replace the Power Management Board
E-18	Problem with DC/DC circuitry	Replace the System/CPU PCA
E-19	Problem with DC/DC circuitry	Replace the System/CPU PCA
E-22	Motor Temperature sensor failedProblem with U37 circuitry	Replace Motor Replace System/CPU PCA
E-23	Internal watchdog on DSP failed	Replace System/CPU PCA



E-68	• Error indicates a short in the Li-Ion battery discharge path on the Power Management PCA. Normally, the Host software recognizes a useable Detachable and Internal battery and attempts to disable the discharge path of the Internal Battery. Since the Internal Battery discharge path has failed shorted, current will be shared by the Internal and Detachable Battery	Replace Power Management PCA
E-78	Unable to read a valid time from the RTC chip	Replace the System/CPU PCA
E-79	The time read from the RTC does not match time on Host CPU	Replace the System/CPU PCA
E-119	Battery charging is detected when there is no AC present	Replace Power Management PCA
E-122	Unable to write to Event Log	Replace the System/CPU PCA
E-123	Error indicates a shorted Q29 in the Li-Ion Battery discharge path on the Power Management PCA. This MOSFET prevents a low mA sneak current from the Internal Battery when the discharge path is disabled by either the Host software or the hardware	Replace the Power Management PCA
E-149	• Before a drift is computed for this sensor, its autonull valve is opened to atmosphere. A test is performed to ensure that the valve has fully opened in accordance with the time provided by the technical specification. This is performed by ensuring that the control pressure has dropped significantly enough after the valve has opened and a de-bounce time has expired. If the pressure reading before the valve was opened was not high enough (due to ventilator therapy settings) to provide a valid criteria regarding a significant drop in pressure, then the check is not performed. When this condition is in effect for a consecutive number of attempted drifts, the error is reported	Replace Sensor Board Replace System/CPU PCA
E-159	Problem with power management board or batteries - unit would not go into Sleep when AC Power removed	Replace Power Management PCA Replace System/CPU PCA
E-198	Circuitry failure	Replace the System/CPU PCA
	I	l .

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 Detachable Battery harness failure Power Management Board Failure Circuitry failure 	 Replace Detachable Battery harness Replace the Power Management PCA Replace the System/CPU PCA
Circuitry failure	Replace the System/CPU PCA
 Power Management Board Failure Circuitry failure 	Replace the Power Management PCA Replace the System/CPU PCA
Circuitry failure	Replace the System/CPU PCA Replace Interface PCA
U3 failureCPLD failure	Replace System/CPU PCA
• Error indicates an open Q13, Q14 in the Lead Acid battery discharge path on the Power Management PCA. When Pb Acid is selected by the hardware or software, VBATTSENSE would be low. If AC is disconnected, there would be no battery backup from the Li-lon battery	Replace Power Management PCA
One or more "Ventilator Service Recommended" errors are active in the system	Examine the Significant Event Log error log. The first column of data contains the first error code that is causing this error message
	Address according to this first error code
	Each time you address an error, cycle the motor (OFF to ON to OFF) and check the Significant Event Log again until this alarm is no longer sounded
	Remove all power from the unit and then reapply power, then turn the motor ON to see if this message is still reported
O ₂ sensor failed	Replace Interface PCA
• Internal O ₂ sensor at end of life or watchdog failure	Replace OBM PCA
 Unit not calibrated O₂ Sensor Calibration Table corrupted 	Recalibrate Replace System/CPU PCA
	Power Management Board Failure Circuitry failure Power Management Board Failure Circuitry failure Circuitry failure Circuitry failure U3 failure CPLD failure Fror indicates an open Q13, Q14 in the Lead Acid battery discharge path on the Power Management PCA. When Pb Acid is selected by the hardware or software, VBATTSENSE would be low. If AC is disconnected, there would be no battery backup from the Li-lon battery One or more "Ventilator Service Recommended" errors are active in the system Og sensor failed Internal Og sensor at end of life or watchdog failure Unit not calibrated



6.7.4 OTHER POWER ALARM ERROR CODES

ERROR CODE	ON-SCREEN MESSAGE	PROBABLE CAUSE	CORRECTIVE ACTION
E-53	High Vte	 The measured Exhaled Tidal Volume is greater than or equal to the alarm setting Flow Sensor problem Active exhalation valve problem 	 Check device and circuit setup if patient circuit is available Check the Active Exhalation Valve functionality, if it is attached Check the alarm settings against the therapy settings. Check the tubing for leaks, kinks, or blockages Replace the Sensor PCA and recalibrate
E-54	Low Vte	 The measured Exhaled Tidal Volume is less than or equal to the alarm setting High leak Flow sensor problem Active exhalation valve problem 	 Check device and circuit setup if patient circuit is available Check the circuit tubing for pinched or blocked tubes if patient circuit is available Check the circuit for leaks if patient circuit is available Check the Active Exhalation Valve functionality, if it is attached Check the alarm settings against the therapy settings Check the tubing for leaks, kinks, or blockages Replace the Sensor PCA and recalibrate
E-81	Card Error	Faulty SD Card	Use different SD Card
E-83	Card Error	 Foreign object inserted in card slot Unformatted card Card prematurely removed 	Use correct MMC/SD card Re-insert card Replace System/CPU PCA



E-84	Card Error	 Foreign object inserted in card slot Unformatted card Card prematurely removed 	 Use correct MMC/SD card Re-insert card Replace System/CPU PCA
E-85	Card Error	 Foreign object inserted in card slot Unformatted card Card prematurely removed 	Use correct MMC/SD card Re-insert card Replace System/CPU PCA
E-86	Card Error	 Foreign object inserted in card slot Unformatted card Card prematurely removed 	Use correct MMC/SD card Re-insert card Replace System/CPU PCA
E-87	Card Error	 Foreign object inserted in card slot Unformatted card Card prematurely removed 	 Use correct MMC/SD card Re-insert card Replace System/CPU PCA
E-88	Card Error	 Foreign object inserted in card slot Unformatted card Card prematurely removed 	Use correct MMC/SD card Re-insert card Replace System/CPU PCA
E-89	Card Error	 Foreign object inserted in card slot Unformatted card Card prematurely removed 	Use correct MMC/SD card Re-insert card Replace System/CPU PCA
E-93	Keypad Stuck	Key is providing reading that it has been held down for 2 minutes	 Check the Keypad for stuck keys Replace the Front Panel PCA Replace the System/CPU PCA



E-109	High Expiratory Pressure	Unit is not reaching the exhaled pressure setting; Patient Pressure during exhalation is greater than or equal to Exhaled Pressure Setting = 5 cm for 5 seconds or more	 Check device and circuit setup if patient circuit available Check the circuit tubing for pinched or blocked tubes if patient circuit available Check the Active Exhalation Valve functionality, if it is attached Check the tubing for leaks, kinks, or blockages Replace the Sensor PCA and recalibrate
E-110	Low Inspiratory Pressure	 Unit is not reaching the inhalation pressure setting; In CPAP, S, S/T, T, PC, and PC-SIMV modes, the lowest Patient Pressure delivered during inhalation is less than the lowest inhalation pressure - 5 cm; In CV, AC, and CIMV modes, the lowest Patient Pressure delivered during inhalation is less than the Low PIP Alarm setting Low pressure sensor reading Large amount of drift Pinched/blocked tubing 	 Check device and circuit setup if patient circuit available Check the circuit tubing for pinched or blocked tubes if patient circuit available Check the circuit for leaks if patient circuit is available Check the Active Exhalation Valve functionality, if it is attached Check the tubing for leaks, kinks, or blockages Replace the Sensor PCA and recalibrate
E-111	High Respiratory Rate	 The measured Breath Rate is greater than or equal to the alarm setting False triggering Alarm/setting mismatch Spontaneous breathing above the alarm 	 Check device and circuit setup if patient circuit available Check the circuit tubing for pinched or blocked tubes if patient circuit available Check the circuit for leaks if patient circuit is available Check the Active Exhalation Valve functionality, if it is attached Check the alarm settings against the therapy settings Check the tubing for leaks, kinks, or blockages Replace the Sensor PCA and recalibrate

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E-112	Low Respiratory Rate	 The measured Breath Rate is less than or equal to the alarm setting Alarm/setting mismatch Spontaneous breathing below the alarm high leak 	 Check device and circuit setup if patient circuit available Check the circuit tubing for pinched or blocked tubes if patient circuit available Check the circuit for leaks if patient circuit is available Check the Active Exhalation Valve functionality, if it is attached Check the alarm settings against the therapy settings Check the tubing for leaks, kinks, or blockages Replace the Sensor PCA and recalibrate
E-113	High Minute Ventilation	 The measured Minute Ventilation is greater than or equal to the alarm setting Alarm/setting mismatch Low breath rate (leak) High exhaled tidal volume (flow sensor problem, active exhalation valve problem) 	 Check device and circuit setup if patient circuit available Check the circuit for leaks if patient circuit is available Check the Active Exhalation Valve functionality, if it is attached Check the alarm settings against the therapy settings Check the tubing for leaks, kinks, or blockages Replace the Sensor PCA and recalibrate



E-114	Low Minute Ventilation	 The measured Minute Ventilation is less than or equal to the alarm setting Alarm/setting mismatch High leak High breath rate Low exhaled tidal volume (flow sensor problem) 	 Check device and circuit setup if patient circuit available Check the circuit tubing for pinched or blocked tubes if patient circuit available Check the circuit for leaks if patient circuit is available Check the Active Exhalation Valve functionality, if it is attached Check the alarm settings against the therapy settings Check the tubing for leaks, kinks, or blockages Replace the Sensor PCA and recalibrate
E-115	High Vti	The measured inhaled tidal volume is greater than or equal to the alarm setting Flow sensor problem Active exhalation valve problem	 Check device and circuit setup if patient circuit available Check the Active Exhalation Valve functionality, if it is attached Check the alarm settings against the therapy settings Check the tubing for leaks, kinks, or blockages Replace the Sensor PCA and recalibrate
E-116	Low Vti	 The measured inhaled tidal volume is less than or equal to the alarm setting High leak Flow sensor problem Active exhalation valve problem 	 Check device and circuit setup if patient circuit available Check the circuit tubing for pinched or blocked tubes if patient circuit available Check the circuit for leaks if patient circuit is available Check the Active Exhalation Valve functionality, if it is attached Check the alarm settings against the therapy settings Check the tubing for leaks, kinks, or blockages Replace the Sensor PCA and recalibrate

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E-118	Low Circuit Leak	 The leak in the system is too small Wrong circuit Blocked tubes Sensor problems 	 Check device and circuit setup Check the circuit tubing for pinched or blocked tubes Check the circuit setting against the circuit being used Check the tubing for leaks, kinks, or blockages Replace the Sensor PCA and recalibrate
E-120	Apnea	 Spontaneous breathing has not been detected within the alarm time High leak 	 Check device and circuit setup Check the circuit tubing for pinched or blocked tubes Check the circuit for leaks Check the Active Exhalation Valve functionality, if it is attached Check the tubing for leaks, kinks, or blockages Replace the Sensor Board PCA and recalibrate
E-121	Circuit Disconnect	 High flow condition has been detected High leak Flow Sensor problem 	 Check device and circuit setup Check the circuit tubing for pinched or blocked tubes Check the circuit for leaks Check the Active Exhalation Valve functionality, if it is attached Check the tubing for leaks, kinks, or blockages Replace the Sensor Board PCA and recalibrate
E-162	Replace Detachable Battery	 Detachable Li-Ion has a TDA due to Discharge Overcurrent or Discharge Short Current PMB fault System Board fault Detachable Li-Ion fault 	 Replace Detachable Li-Ion Replace Power Management PCA Replace System CPU PCA



E-165	Low Expiratory	When the Expiratory pressure is less than or equal to the EPAP/CPAP/PEEP - 5 pressure units in value for 5 seconds or more Deactivated if the Attained Expiratory Pressure > EPAP/PEEP/CPAP - 5 pressure units	 Check the circuit tubing for pinched or blocked tubes if patient circuit available Check the circuit for leaks if patient circuit available Check the Active Exhalation Valve functionality, if it is attached Check the tubing for leaks, kinks, or blockages Replace the sensor board
E-170	High Inspiratory Pressure	 Unit is exceeding the inhalation pressure setting; IN CPAP, S, S/T, T, PC, and PC-SIMV modes, the highest Patient Pressure delivered during inhalation is greater than the highest inhalation pressure + 5 cm; In CV, AC, and SIMV modes, the highest Patient Pressure delivered during inhalation is greater than the High PIP Alarm setting High pressure Sensor reading Large amount of drift Pinched/blocked tubing 	 Check the circuit tubing for pinched or blocked tubes if patient circuit available Check the circuit for leaks if patient circuit available Check the Active Exhalation Valve functionality, if it is attached Check the tubing for leaks, kinks, or blockages Replace the Sensor PCA
E-171	High Inspiratory Pressure	 Unit is exceeding the inhalation pressure setting; IN CPAP, S, S/T, T, PC, and PC-SIMV modes, the highest Patient Pressure delivered during inhalation is greater than the highest inhalation pressure + 5 cm; In CV, AC, and SIMV modes, the highest Patient Pressure delivered during inhalation is greater than the High PIP Alarm setting High pressure sensor reading Large amount of drift Pinched/blocked tubing 	 Check the circuit tubing for pinched of blocked tubes if patient circuit is available Check the circuit for leaks if patient circuit is available Check the Active Exhalation Valve functionality, if it is attached Check the tubing for leaks, kinks, or blockages Replace the Sensor PCA

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_	T _	T	
E-173	Start On Battery	Vent was started on battery power	 None - recorded for informational purposes to indicate the unit was started without AC power.
E-176	Replace Detachable Battery	 Can't communicate with Detachable Li-Ion battery and charger is not trying to wake-up charge the battery. Could take up to 210 seconds for alarm to be reported because the charger is trying to wake-up charge the Detachable Li-Ion PMB fault Detachable Li-Ion fault 	 Replace the Power Management PCA Replace the Detachable Li-Ion
E-178	Replace Detachable Battery	 Detachable Li-Ion battery not authentic Fake battery PMB fault Detachable Li-Ion fault 	Replace Detachable Li-Ion Replace Power Management PCA
E-179	AC Power Disconnected	 AC was disconnected PMB fault AC Power Supply fault Power Cord fault A/D channel fault 	 Verify AC connected Replace Power Management PCA Replace AC Power Supply Replace System/CPU PCA
E-180	External Batt Disconnected	 Lead Acid was disconnected Power Cord fault PMB fault A/D channel fault 	 Connect Lead Acid Verify Lead Acid Battery Cord Replace Power Management PCA Replace System/CPU PCA
E-181	Detach Batt Disconnected	 Detachable Li-Ion was disconnected Li-Ion connector fault PMB fault A/D channel fault 	 Verify Detachable Li-Ion connected Replace Detachable Li-Ion cable Replace Power Management PCA Replace System/CPU PCA
E-183	Low External Battery	Low battery - Lead Acid has 20 minutes run time remaining and it is last available power source 	Charge Lead Acid Battery Replace Lead Acid Battery



E-184	Low Detachable Battery	• Low battery - Detachable Lilon lon has ≤ 20 minutes run time remaining and it is last available power source • Charge Detachable Li-lon Battery • Replace Detachable Li-lon Battery
E-185	Low Internal Battery	• Low battery - Internal Li-Ion has ≤ 20 minutes run time remaining and it is last available power source
E-186	Low External Battery	 Depleted battery - Lead Acid has ≤ 10 minutes run time remaining and it is last available power source Charge Lead Acid battery Replace Lead Acid battery
E-187	Low Detachable Battery	• Depleted battery - Detachable Li-lon has ≤ 10 minutes run time remaining and it is last available power source • Charge Detachable Li-lon Battery • Replace Detachable Li-lon battery
E-188	Low Detachable Battery	Depleted battery - Internal Lilon has ≤ 10 minutes run time remaining and it is last available power source Charge Internal Li-Ion Battery Charge Internal Li-Ion Battery
E-192	Replace Detachable Battery	 AFE chip does not receive the appropriate frequency on the WDI pin from the Gas Gauge chip. Both of these chips are in the Li-lon battery pack Replace Detachable Li-lon battery Replace Power Management Board Detachable Li-lon fault
E-194	Replace Detachable Battery	Detachable Li-lon Battery Permanent Failure Battery battery
E-196	Replace Detachable Battery	 Detachable Li-Ion State of Health ≤ 50%. Full Charge Capacity (FCC) is less than 51% of the design capacity Detachable Li-Ion FCC error Detachable Li-Ion fault Detachable Li-Ion end of life Cycle the Detachable Li-Ion (discharge battery so that capacity change is > 50%, then rest for > 2 hours. Charge to 100%, then rest for > 2 hours) Replace Detachable Li-Ion battery
E-206	Replace Detachable Battery	 Corruption of battery gas gauge Cycle the Detachable Li-lon (diecharge battery so that capacity change is > 50%, then rest for > 2 hours. Charge to 100%, then rest for > 2 hours)

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E-209	Replace Detachable Battery	 Detachable Li-Ion Cycle Count ≥ 5500 cycles Detachable Li-Ion end of life 	Replace Detachable Li-Ion Battery
E-218	Batt Discharge Stopped-Temp.	Internal Li-lon charge current greater than 4 Amps for greater than 2 sec. Battery charge FET is turned off. High temperature during discharge. Battery recovers if temperature is less than 55° C.	 Inspect fans Replace cooling fan(s)
E-219	Batt Discharge Stopped-Temp.	Detachable Li-lon charge current greater than 4 Amps for greater than 2 sec. Battery charge FET is turned off. High temperature during discharge. Battery recovers if temperature is less than 55° C.	 Inspect fans Replace cooling fan(s)
E-230	Battery Not Charging - Temp.	 Internal Li-Ion temp > 45° C or < 0° C when charging is initiated. Resets when temp ≤ 42° C and ≥ 3° C Internal Li-Ion temp > 50° C or < -2.5° C during charging. Resets when temp 42° C and ≥ 0.5° C High/low ambient temperature Cooling fan(s) fault Detachable Li-Ion fault 	 Inspect fans Replace cooling fan(s) Replace Internal Li-Ion
E-231	Battery Not Charging - Temp.	 Detachable Li-lon temp > 45° C or < 0° C when charging is initiated. Resets when temp ≤ 42° C and ≥ 3° C Detachable Li-lon temp > 50° C or < -2.5° C during charging. Resets when temp 42° C and ≥ 0.5° C High/low ambient temperature Cooling fan(s) fault Detachable Li-lon fault 	 Inspect fans Replace cooling fan(s) Replace Detachable Li-Ion



E-234	Upgrade Failed Screen	Unable to open the new software file on the card	 Reformat card and replace new software file on the card Re-insert card, retry upgrade
E-235	Upgrade Failed Screen	 Unable to read the new software file on card The new software file on the card is corrupt 	 Reformat card and replace new software file on the card Re-insert card, retry upgrade
E-236	Upgrade Failed Screen	The user tried to upgrade to an older version of software	 Reformat card and replace new software file on the card Re-insert card, retry upgrade
E-237	Upgrade Failed Screen	The user tried to upgrade to a version of software that is not intended for the Trilogy Ventilator	 Reformat card and replace new software file on the card Re-insert card, retry upgrade
E-238	Upgrade Failed Screen	The new software file on the card is corrupt	 Reformat card and replace new software file on the card Re-insert card, retry upgrade
E-239	Upgrade Failed Screen	Unable to program the new software on the Host CPU	Replace System/CPU PCA
E-240	Upgrade Failed Screen	The user tried to upgrade to a version of software that is not intended for the Trilogy Ventilator	 Reformat card and replace new software file on the card Re-insert card, retry upgrade
E-241	Upgrade Failed Screen	Unable to program the new software on the DSP	Replace System/CPU PCA
E-242	Upgrade Failed Screen	Unable to program the new software on the Host CPU	 Reformat card and replace new software file on the card. Re-insert card, retry upgrade Replace System/CPU PCA
E-245	Software stops blower and Loss of All Power sounds	Last battery depleted	Replace or charge the battery
E-246	Internal Battery Not Charging	 Unable to charge Internal Lilon battery after 30 minutes PMB fault Internal Li-lon fault Detachable Li-lon fault 	 Replace Internal Li-Ion Replace Detachable li-Ion Replace Power Management PCA

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E-247	Detach Battery Not Charging	 Unable to charge Detachable Li-lon battery after 30 minutes PMB fault Detachable Li-lon fault Internal Li-lon fault 	 Replace Detachable Li-Ion Replace Internal Li-Ion Replace Power Management PCA
E-266	Replace Detachable Battery	Checks if Li-lon battery is present by attempting to communicate when the Charger Chip safety signal indicates the battery is not present. Safety signal from the Li-lon may be disconnected or Charger Chip may have fault so it is not detecting a Li-lon battery	 Replace the Power Management PCA Replace Detachable Battery
E-280	External Battery Depleted	Lead Acid was depleted	Replace or charge Lead Acid
E-281	Detachable Battery Depleted	Detachable Li-lon was depleted	Replace or Charge Detachable Li-lon
E-282	Internal Battery Depleted	 Internal Li-Ion was depleted Internal Li-Ion discharge FET off 	Charge Internal Li-Ion Replace Internal Li-Ion
E-283	Upgrade Screen Complete	Software upgrade completed successfully	None - recorded for informational purposes to indicate that a software upgrade was completed
E-291	Check Circuit	 Patient circuit does not match Circuit Type Tubing on active circuit not connected properly Tubing on active circuit fell off One of dual sensors failed 	 Change circuit or Circuit Type Reconnect tubing correctly Reconnect tubing Install correct porting block Replace Sensor PCA
E-299	Card Error	When the SD Card does not have enough memory available to write EDF (3 days of Waveform and 1 year of Annotations and Detailed data) and Event Log (CSV) Files	 Erase unnecessary files on card. Ensure card > 75 MB Use new card



E-300	Card Error	• When the Cord is detected as	• Chook write protect ewitch an
<i>E-300</i>	Card Entir	When the Card is detected as a write protected SD Card	 Check write protect switch on card Use new card Replace System/CPU PCA
E-301	Card Error	Error in creating files/ directories on the SD Card	Use new card Replace System/CPU PCA
E-305	High Internal Oxygen	 Disconnected O₂ tubing O₂ level between 25% and 35% 	 Connect/replace O₂ tubing Replace Interface PCA
E-307	High Temperature	It could be caused by air stream temperature, motor temperature or batteries temperature	 Check/replace fans Replace batteries Replace motor Replace air stream temperature sensor
E-308	High Temperature	It could be caused by air stream temperature, motor temperature or batteries temperature	 Check/replace fans Replace batteries Replace motor Replace air stream temperature sensor
E-309	Check External Battery	 Lead Acid is useable and current is being drawn from the Li-lon battery External Battery Cable fault Faulty connection to Leak Acid battery terminals PMB fault A/D channel fault High impedance Lead Acid battery 	 Check connections Replace External Battery Cable Replace Lead Acid Battery Replace Power Management PCA Replace System/CPU PCA
E-312	Prescription Change screen displayed at time of Rx Update	If usable Rx found on the SD Card for the device.	None - recorded for informational purposes that the prescription was read successfully from the SD card.
E-313	Prescription Review screen displayed at time of Rx Update	Rx from the SD Card is ready for user review	None - recorded for informational purposes that the prescription was read successfully from the SD card.

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E-314	Failed -Card is read only	If the Rx is only for the current device and the card is read only	Check write protect switch on card Replace System/CPU PCA
E-315	Failed - Serial Number	If the Serial Number in the Rx Card does not match the Serial Number of the device	None - recorded for informational purposes that the serial number in the prescription does not match the serial number for the device.
E-316	Failed - Circuit Type	If the circuit type in the Rx on the card does not match the Circuit Type in the device	 None - recorded for informational purposes that the circuit type in the prescription does not match the circuit type for the device.
E-317	Prescription Change Failed	Failure to update the Rx due to the errors in the Rx settings	None - recorded for informational purposes to indicate that the prescription had bad settings.
E-318	Prescription Change Complete	Rx update is complete	None - recorded for informational purposes to indicate that the prescription update was completed successfully.
E-319	Prescription Change Cancelled	Rx update is cancelled by the user by either UI option or by pulling out of the card before Rx Update is complete	None - recorded for informational purposes indicate that the prescription update was not completed.
E-320	Prescription Change Failed	failure to update the Rx due to error in the file format or errors in reading the file	None - recorded for informational purposes to indicate that the prescription data on the card could not be read.
E-321	Prescription Change failed	Rx version incorrect for the device	None - recorded for informational purposes to indicate that the version of the prescription data on the SD card was bad.
E-322	Prescription Change Failed	Rx checksum is incorrect	None - recorded for informational purposes to indicate that the prescription on the SD card had a bad checksum.



E-339	Pressure Sensor	Temperature Table used in	Run unit to heat up, turn unit off
_ 333	Drift - turn off for 20 secs.	temperature rable used in temperature compensation for dP sensor is not sufficient for use	(leave plugged in), let unit cool down to get more data in Temperature Table
			Replace Sensor Board
E-342	High Priority	Pressure at Oxygen Blending Module inlet below 40 psi	 Check supply pressure Check inlet pressure sensor connections Replace OBM PCA
			Replace OBIVI FCA
E-343	High Priority	Pressure at Oxygen Blending Module above 87 psi	 Check supply pressure Check inlet pressure sensor connections Replace OBM PCA
E-348	Upgrade Failed Screen	OBM did not accept new software version	 Reformat card and replace new software file on card Re-insert card; retry upgrade
E-351	Low Oxygen Flow	• Delivered oxygen concentration less than FiO ₂ set point - 10%	 Check supply pressure Check inlet pressure sensor connections Check internal flow element for leaks Replace OBM Module Replace OBM PCA
E-352	High Oxygen Flow	Delivered oxygen concentration greater than FiO ₂ setpoint + 10%	 Check inlet pressure Replace OBM valve Replace OBM PCA
E-359	High Internal Oxygen	O ₂ level inside OBM > 30%	 Connect O₂ tubing in OBM Disconnect O₂ source or replace sensor in OBM
E-360	Low Vte	 The desired tidal volume cannot be delivered within the limits of the IPAP MIn and Max settings High leak Flow Sensor problem 	 Check the tubing (inside and outside unit) for leaks, kinks, or blockages Replace the sensor board and recalibrate

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CHAPTER 7: MAINTENANCE

7.0 CHAPTER OVERVIEW

This chapter describes the Maintenance intervals and procedures for the Trilogy Ventilator.

7.1 ROUTINE MAINTENANCE

Routine maintenance involves periodic checking, cleaning, and or replacing the following item as necessary:

- Air Inlet Filter
- Enclosures

7.1.1 AIR INLET FILTER

Under normal usage, you should clean the inlet filter at least once every two weeks and replace it with a new filter every six months or sooner if needed.

7.1.2 ENCLOSURES

Commensurate with hospital or homecare policies, Respironics recommends cleaning the cabinet and inspecting for damage as necessary.

7.2 PREVENTIVE MAINTENANCE

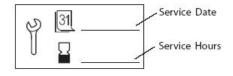
The following Preventive Service procedures will be followed by authorized service personnel at the intervals outlined in the maintenance schedule below. Preventive Maintenance must be performed by Respironics or a Respironics' trained location.

Your new Philips Respironics Trilogy ventilator includes a PM Service Due label affixed to the bottom of your device. The Service Date line on the label is blank, which allows you to document the date when service is due. The Service Hours line will have 10000 hours recorded. Service is due every 10000 blower hours or 24 months whichever comes first depending on the usage of the device or 17,500 blower hours for blower maintenance.

During the initial install, you will need to record the date and hours (if not already written) on the blank label to maximize your in-service interval. Record the service date as 24 months from the date first placed on a patient. Record the hours as 10000 blower hours for initial usage.

The device blower hours are listed in the device software in the information menu.

A new blank label should be applied to the Trilogy Ventilator after the 10000 hours or 24 months maintenance and after the 17,500 Blower Maintenance has been performed. The label should be completed according to the table below with the new next maintenance due blower hours and new next maintenance due date (should be 24 months from date of completed Maintenance) before returning the ventilator.



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WHEN TO COMPLETE	LABEL INFORMATION	LABEL COMPLETED BY
First ever product use	Record 10000 on the bottom line and 24 months from date of install on the top line.	Trilogy Dealer
After first 10000 hours / 24 months maintenance completion	Record 17,500 on the bottom line and add date 24 months from 10000 / 24 months maintenance completion date. For example if the maintenance was completed on 01/01/2010 (MM/DD/YYYY) the date recorded on the top line should be 1/1/2012 (MM/DD/YYYY).	Service Technician
After first 17,500 blower maintenance completion	Record 27,500 on the bottom line and add date 24 months from blower maintenance completion date. For example if the maintenance was completed on 01/01/2010 (MM/DD/YYYY) the date recorded on the top line should be 01/01/2012 (MM/DD/YYYY).	Service Technician
After second 10000 hours / 24 months maintenance completion	Record 35,000 on the bottom line and add date 24 months from 10000 hours / 24 months maintenance completion date. For example if the maintenance was completed on 01/01/2010 (MM/DD/YYYY) the date recorded on the top line should be 01/01/2012 (MM/DD/YYYY).	Service Technician
After second 17,500 blower maintenance completion	Record 45,000 on the bottom line and add date 24 months from blower maintenance completion date. For example if the maintenance was completed on 01/01/2010 (MM/DD/YYYY) the date recorded on the top line should be 01/01/2012 (MM/DD/YYYY).	Service Technician



NOTES

NOTE 1: The hours used above may vary based on the return. If the device is returned for blower maintenance and there were 19,000 blower hours recorded the next maintenance due hours should be recorded as 29,000.

NOTE 2: After the initial 10000 hours or 24 months Maintenance has been performed, if a device is returned the second time using the date interval and it has not reached the blower maintenance interval perform the 10000 hours / 24 months maintenance again and the label should be completed to show the due date for the Blower Maintenance again.

For example - Service completes the 10000 hours / 24 months Maintenance on 01/01/2010 (MM/DD/YYYY), they complete the Preventive Maintenance to show 17,500 as the Maintenance due hours and 1/1/2012 for the Maintenance due date. On 01/01/2012 (MM/DD/YYYY) the device was returned for Maintenance as the label states and there are only 12,000 hours recorded on the blower. At this point you may contact the customer to see if they would like the blower maintenance performed early.

If they decide to have the blower maintenance performed complete the maintenance and record 22,000 on the bottom line for the next maintenance due hours and 1/1/2014 on the top line for the next maintenance due date.

If they decide to not have the blower maintenance performed perform the 10000 hours / 24 months maintenance and once completed record 17,500 on the bottom line for next maintenance due hours and 1/1/2014 on the top line for the next maintenance due date.

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7.2.1 TRILOGY RECOMMENDED MAINTENANCE SCHEDULE

TIME / HOURS OF SERVICE	RECOMMENDED MAINTENANCE
Prior to initial use on a patient	 Charge Internal and Detachable Batteries to 100% capacity by plugging ventilator into an AC power source for up to 8 hours
	 Perform System Checkout Procedure per Trilogy Clinical Manual
Prior to long term storage after initial use	 Charge both Internal and Detachable batteries to 100% capacity prior to storage.
While in storage after initial connection to AC Power, every 3 months	 Recharge internal and detachable batteries to 100% capacity by plugging ventilator into an AC power source. Batteries should recharge in 8 hours or less.
If in use, every two weeks	Inspect and clean air inlet filter
If in use, every 6 months	Replace air inlet filter
	 Inspect enclosures and external connections for damage and contact Philips Respironics service if necessary
	 Inspect power cord for damage and replace if necessary.
Every 10000 hours or 24 Months, whichever comes first	Refer to appropriate section below.
Every 17,500 hours	Refer to appropriate section below.



7.2.2 10000 Hours or 24 Months (Whichever Comes First) Maintenance Items

ITEM	ACTION
Blower Hours/Date of Maintenance	Record the blower hours and maintenance date.
Lithium Ion Battery Cycle Count Limit	Check the Cycle Count on the Detachable and Internal Batteries. Procedure listed below.
	The service technician has the ability to access the number of cycles that both the Detachable and Internal batteries have gone through. The Internal Battery has the capacity to reach 475 cycles and the Detachable Battery has the capacity to reach 500 cycles before the battery life is exhausted. Respironics recommends checking the cycle count of both batteries during the annual preventive maintenance. By viewing and recording the cycle count the service technician will be able to determine if the batteries need to be replaced based on the number of cycles accumulated since the last annual preventive maintenance that was performed on the Trilogy. Below is an example of how the technician may determine if the batteries need replaced.
	Internal Battery Cycle Capacity: 475 cycles Internal Battery Cycles Used: 125 cycles Internal Battery Cycles remaining: 350 cycles
	Detachable Battery Cycle Capacity: 500 cycles Detachable Battery Cycles Used: 125 cycles Detachable Battery Cycles remaining: 375 cycles
	This example shows that the Trilogy Internal Battery and Detachable Battery went through 125 cycles from the last annual preventive maintenance leaving 375 cycles for the Detachable Battery and 350 cycles for the Internal Battery. By using this method, the service technician can assume that the batteries will have enough cycles remaining until the next annual preventive maintenance is performed.
	Respironics recommends that you replace the Detachable and Internal batteries when they have logged 400 or greater cycles.
Replace Inlet Filter (RI p/n 1035443)	Replace the Inlet Filter.
Outer Enclosure	Clean the Outer Enclosure.
SD Card Interface	Inspect the Door Hinge and foam. Inspect the Latch/Release mechanism.
Connectors	Verify the integrity of connectors on the rear of the unit, including the AC cord retainer.



İTEM	ACTION
Porting Blocks	Check O-Rings around the porting blocks for cracks, wear, or incorrect fit. Lubricate or replace the porting block as necessary.
Detachable Battery (if present)	Verify the latch and guide mechanism is intact and functional. Compare the capacity LEDs on the Detachable Battery Pack to the Detachable Battery Icon on the User Interface. The Charge levels should be within 1 level of each other.
Handle	Verify the Handle is free of defects. Check O-Rings around the handle for cracks, wear, or incorrect fit. Lubricate or replace as necessary.
Rubber Feet	Verify that all of the Rubber Feet are present.
Power Cord	Inspect the power Cord for any cracks/cuts.
External Tubing	Clean or replace all external tubing as necessary.
Cables and Internal Tubing	Remove the Bottom Enclosure and verify the integrity of cable connectors and internal tubing.
Cleaning	Remove the Bottom Enclosure and vacuum out the bottom of the unit using an Anti-static vacuum.
Perform Field Service Application (FSA) Post Test Procedure or Performance Verification (PV) Tool	Preform the Field Service Application Post Test Procedure ^a or the Performance Verification Tool ^b .

^a If the unit fails the FSA Post Test Procedure refer to the Troubleshooting Chapter of this service manual to determine the corrective action that needs taken. Once the corrective action has been performed run the complete FSA.

^b If the unit fails the PV tool, return the device to an authorized Service Facility for troubleshooting and/or repair. Once the corrective action has been performed run the complete FSA.



7.2.3 17, 500 BLOWER MAINTENANCE ITEMS

ITEM	Action
Blower Hours/Date of Maintenance	Record the blower hours and maintenance date.
Blower Assembly (RI p/n 1054951)	Replace the Blower Assembly with a new Blower Assembly.
Lithium Ion Battery Cycle Count Limit	Check the Cycle Count on the Detachable and Internal Batteries. Procedure listed below.
	The service technician has the ability to access the number of cycles that both the Detachable and Internal batteries have gone through. The Internal Battery has the capacity to reach 475 cycles and the Detachable Battery has the capacity to reach 500 cycles before the battery life is exhausted. Respironics recommends checking the cycle count of both batteries during the annual preventive maintenance. By viewing and recording the cycle count the service technician will be able to determine if the batteries need to be replaced based on the number of cycles accumulated since the last annual preventive maintenance that was performed on the Trilogy. Below is an example of how the technician may determine if the batteries need replaced.
	Internal Battery Cycle Capacity: 475 cycles Internal Battery Cycles Used: 125 cycles Internal Battery Cycles remaining: 350 cycles
	Detachable Battery Cycle Capacity: 500 cycles Detachable Battery Cycles Used: 125 cycles Detachable Battery Cycles remaining: 375 cycles
	This example shows that the Trilogy Internal Battery and Detachable Battery went through 125 cycles from the last annual preventive maintenance leaving 375 cycles for the Detachable Battery and 350 cycles for the Internal Battery. By using this method, the service technician can assume that the batteries will have enough cycles remaining until the next annual preventive maintenance is performed.
	Respironics recommends that you replace the Detachable and Internal batteries when they have logged 400 or greater cycles.
Replace Inlet Filter (RI p/n 1035443)	Replace the Inlet Filter.
Outer Enclosure	Clean the Outer Enclosure.
SD Card Interface	Inspect the Door Hinge and foam. Inspect the Latch/Release mechanism.



İTEM	ACTION
Connectors	Verify the integrity of connectors on the rear of the unit, including the AC cord retainer.
Porting Blocks	Check O-Rings around the porting blocks for cracks, wear, or incorrect fit. Lubricate or replace the porting block as necessary.
Detachable Battery (if present)	Verify the latch and guide mechanism is intact and functional. Compare the capacity LEDs on the Detachable Battery Pack to the Detachable Battery Icon on the User Interface. The Charge levels should be within 1 level of each other.
Handle	Verify the Handle is free of defects. Check O-Rings around the handle for cracks, wear, or incorrect fit. Lubricate or replace as necessary.
Rubber Feet	Verify that all of the Rubber Feet are present.
Power Cord	Inspect the power Cord for any cracks/cuts.
External Tubing	Clean or replace all external tubing as necessary.
Cables and Internal Tubing	Remove the Bottom Enclosure and verify the integrity of cable connectors and internal tubing.
Cleaning	Remove the Bottom Enclosure and vacuum out the bottom of the unit using an Anti-static vacuum.
Perform the Complete Field Service (FSA) Application (Pre and Post Test)*	Once the Blower Assembly has been replaced preform the Field Service Application Pre and Post Test Procedures*.

^{*} If the unit fails the Field Service Application Pre and/or Post Test refer to the Troubleshooting section of this service manual to determine the corrective action that needs taken. Once the corrective action has been performed run the complete FSA.

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7.3 TRILOGY VENTILATOR MAINTENANCE RECORD

MODEL NUMBER	SERIAL NUMBER
DATE PURCHASED	BLOWER HOURS
BLOWER ASSEMBLY (Date Replaced)	INLET FILTER (Date Replaced)
INTERNAL BATTERY	DETACHABLE BATTERY
CYCLE COUNT LIMIT (Cycles Remaining)	CYCLE COUNT LIMIT (Cycles Remaining)



7.4 Performance Verification (PV) Tool

The Performance Verification (PV) Tool software will provide the trained user the ability to verify the essential performance and safety of the Trilogy unit under test (UUT).

7.4.1 DOWNLOADING THE PV TOOL

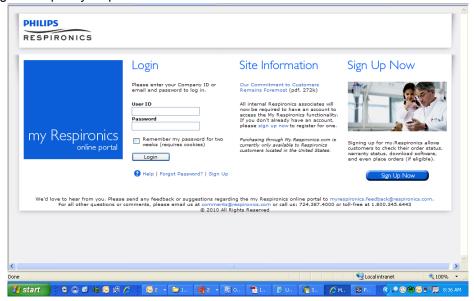
NOTE

Respironics service software is now available at http://my.respironics.com. In the event that you are unable to access this site, log onto http://servicesoftware.respironics.com. to download Respironics service software.

You must be a registered user to download the PV Tool. To become a registered user, you must successfully complete the Trilogy Service Training class and the registration process listed on the website.

Once you have access to download the software, perform the following:

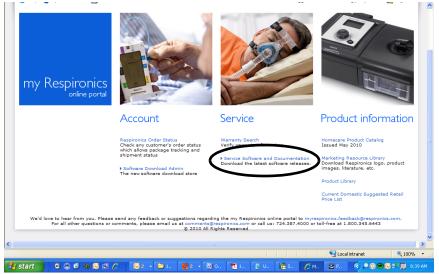
1. Log into http://my.respironics.com.



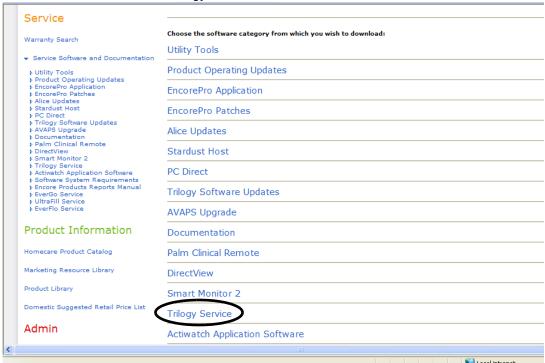
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2. Click on the Service Software and Documentation link.



3. In the menu below select Trilogy Service.



- 4. Click on the Download button adjacent to the PV Tool software.
- 5. When you click on the Download button, the "Run or Save?" window will appear.
- 6. Click on Save to download the software and save it to a specific location on your PC.
- 7. Follow the on-screen prompts to "Save" the software.
- 8. After download is complete, open folder where file was saved.
- 9. Double click the install file to install the application onto your computer.
- 10. Follow the on-screen prompts to complete the installation.

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11. You must restart your computer after this installation. Once the computer restarts there will be a new icon on the desktop screen of the computer.

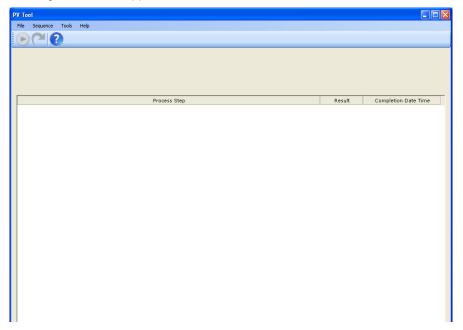
7.4.2 OPENING THE PV TOOL

This section describes how to access the PV Tool after it is installed on your PC along with detailing the Menus and Options that the PV Tool offers.

1. To access the PV Tool, either select the icon from the desktop if you have created one, or select "Start --> All Programs --> Respironics --> Performance Verification Tool".



2. The following screen will appear.



7.4.3 GETTING STARTED

After the user double-click on the Performance Verification Tool desktop icon or selects the Performance Verification Tool option from the Windows Start Menu, the main window appears. The Main window consists of:

- Menu Bar
- Tool Bar
- Overview Panel

MENU BAR

The Menu bar at the top of the screen contains four items:

- File
- Sequence
- Tools
- Help

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File Pull-Down Menu

The file pull- down menu contains the following items:

MENU ITEM	DESCRIPTION
New	Allows the user to begin a new test. The user should select the product they would like to test and then select the open button to open the test program sequence.
Open	Allows the user to access completed tests.
Print	Allows the user to print completed test data sheets. The print option is only available for completed tests.
Exit	Allows the user to exit the PV Tool Program.

Sequence Pull-Down Menu

The sequence pull-down menu contains the following items:

MENU ITEM	DESCRIPTION
Run	Allows the user to begin a test. The user may also use the Run Icon instead of this menu option.
Resume	Allows the user to resume a paused test. The user may also use the Resume Icon instead of this menu option

Tools Pull-Down Menu

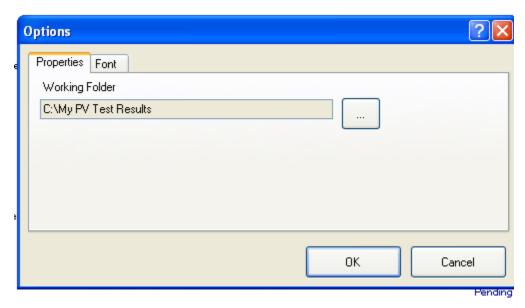
The tools pull-down menu contains the following items:

MENU ITEM	DESCRIPTION
Options	Displays an Options screen, which allows the user to modify his/her working folder or modify font type and size.
RASP	The reset device function allows the user to return the UUT to its original mode in the case that there was an interruption or hang up in the previous test which left the UUT in a non-responsive state.



Options Window

The options pop-up window is show below.



Changing the Properties

Under the Properties tab, the user can change the folder location where the saved test files are stored.

- 1. To change the Working Folder where the test files are saved, click on the button to the right of the field to browse to a new location.
- 2. Select the new folder where the user wants the files to be saved, and click the OK button to return to the Properties tab. The user must select a valid location in the file system.

Changing Font

Under the Font tab, the user can select the font and size the user would like the PV Tool to display.

Help Pull-Down Menu

The help pull-down menu contains the items:

MENU ITEM	DESCRIPTION
PV Tool Help	Displays the Performance Verification instructions for use.
About	Displays the name and software version of the application, as well as copyright, legal, and licensing notices.

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TOOL BAR

The options on the tool bar are enabled or disabled on the current test state.

İTEM	DESCRIPTION
D	Select this icon to run the test.
	Select this icon to resume a test.
?	Select this icon to read attached help files.
	Select this icon to pause a test.
8	Select this icon to reject a test.

OVERVIEW PANEL

The Overview Panel on the Main window displays the test sequence along with session information once a test has been selected.

7.4.4 RUNNING THE TRILOGY PV TOOL

REQUIRED EQUIPMENT

- Computer (Minimum requirements):
 - 400MHZ Processor
 - 256MB RAM
 - 500MB Hard Disk Space
 - CDROM Drive
 - RS-232 Port or USB Port with RS-232 Adapter
 - Windows Pointing Device (Mouse)
 - Video,16-bit color, 800 x 600 Resolution
 - Monitor capable of displaying 16-bit color, 800 x 600 resolution
 - Printer (Optional)



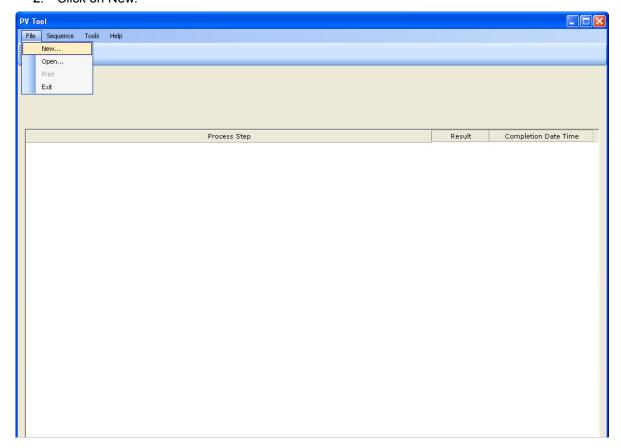
- Operating Systems supported: Windows XP (SP3), Windows Vista (32 and 64 bit), Windows 7 (32 and 64 bit)
- Flow Measurement Device
 - Recommended TSI Certifier FA Plus Test System (Includes tubing, additional pick-off port, and filter) Respironics Part Number 1040310 or any device that meets the following requirements: Capable of measuring Flow 0 200 SLPM in external patient tubing with an accuracy of +/- 2% of reading or 0.1slpm, whichever is greater. Flow Measurement Device must display readings in SLPM to a minimum resolution of one decimal point.
- Pressure Measurement Device
 - Recommended TSI Certifier FA Plus Test System (Includes tubing, additional pick-off port, and filter) Respironics PN 1040310 or any device that meets the following requirements:
 Capable of measuring pressure 0 80 cmH₂O in external patient tubing with an accuracy of +/- 0.3 cmH₂O. Pressure Measurement Device must display readings in cmH₂O to a minimum resolution of one decimal point.
- Atmospheric Pressure Measurement Device
 - Recommended TSI Certifier FA Plus Test System (Includes tubing, additional pick-off port, and filter) Respironics PN 1040310 or any device that meets the following requirements:
 Capable of measuring atmospheric pressure 374.9 890 mmHg (14.76 35 in Hg) with an accuracy of +/- 7.62 mmHg (0.3 in Hg). Atmospheric Pressure Measurement Device must display atmospheric pressure readings in mmHg with a minimum resolution of two decimal points.
- Temperature Measurement Device
 - Recommended TSI Certifier FA Plus Test System (Includes tubing, additional pick-off port, and filter) Respironics PN 1040310 or any device that meets the following requirements:
 Capable of measuring air temperature 0 50 degrees C in external patient tubing with an accuracy of +/- 1 degree C. Temperature Measurement Device must display air temperature in C with a minimum resolution of one degree C.
- Resistance Measurement Device
 - Recommended Fluke 87 V (Includes test leads) Respironics PN 1071681 or any device that meets the following requirements: Capable of measuring resistance 0 – 10 Mohms with an accuracy of 1% of measured range. Resistance Measurement Device must display resistance measurement in ohms with a minimum resolution of one decimal point.
- Additional Required Test Equipment when using Recommended Measurement Devices
 - Trilogy Nurse Call Adapter Cable (Respironics Part Number 1045290)
 - Trilogy Device to PC Data Cable (Respironics Part Number 1046972)
 - SD Card (for loading Trilogy software) or any 1 or 2 Gig SD Card (Respironics Part Number 1051801)
 - Test Orifice, .25" ID (Respironics Part Number 332353)
 - Outlet Port (Respironics Part Number 312710)
 - Whisper Swivel II (Respironics Part Number 332113)
 - Exhalation Porting Block, Active w/PAP (Respironics Part Number 1054670)
 - Smoothbore Tubing; 6ft (Respironics Part Number 622038)



- 3/16" ID Tubing connect Prox pressure to outlet port (Local Supplier or Respironics Part Number 1060747 (Test hardware Kit))
- 1/8" ID Tubing to connect circuit pressure to pressure measurement device (Local Supplier or Respironics Part Number 1060747 (Test hardware Kit))
- Auto DC Battery Cable (Respironics Part Number 1067424)
- Trilogy Detachable Battery (Respironics Part Number 1043570)
- Portable Battery Pack (14.4AH) (Respironics Part Number 1028869) or Deep-Cycle Marine Battery (Local Supplier) or DC Power Source capable of providing 15 VDC and 24 Amps DC (Respironics Part Number 1071678)

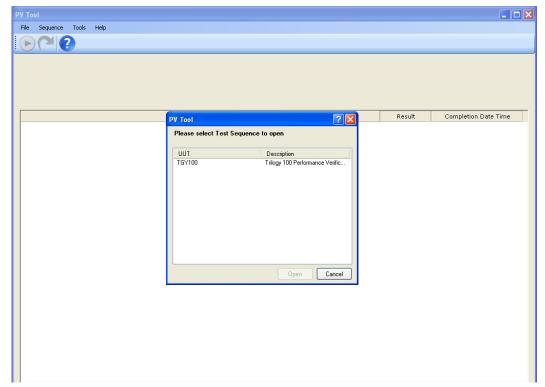
PROCEDURE

- 1. From the main menu, click on File.
- 2. Click on New.

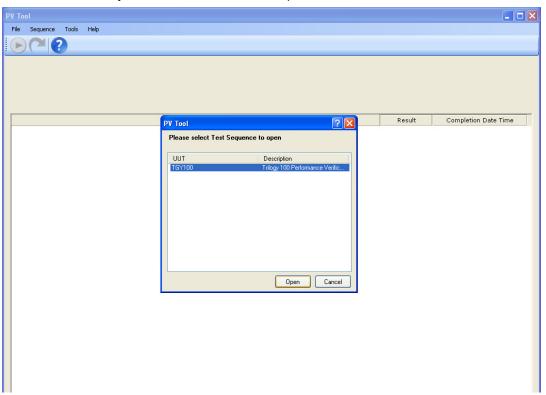




3. The following screen will appear.



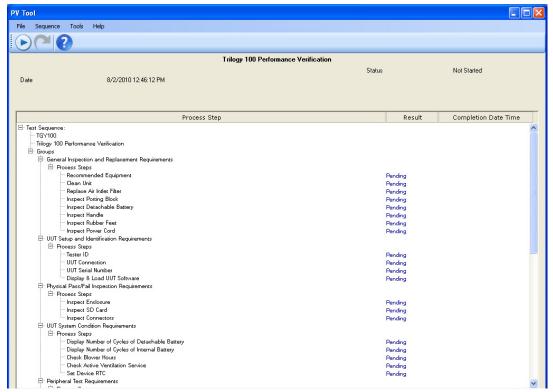
4. Click on the test you want to run and select Open.



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5. The following screen will appear.



- 6. To begin the test either select run from the file menu or click the run icon.
- 7. Using the on-screen prompts and associated help files complete the PV.
- 8. At the end of a successful or failed test a test report will be generated for you to sign and keep with your records. A soft copy of the test report will also be saved to your hard drive.



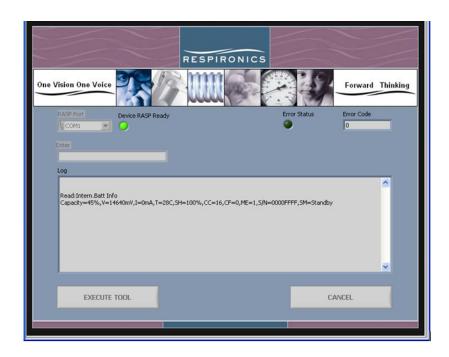
7.5 CHECKING THE INTERNAL BATTERY AND DETACHABLE BATTERY CYCLE COUNTS

1. Double click on the Trilogy Tool Box Icon.

NOTE

If the Trilogy Toolbox software is not downloaded to your computer, refer to the Testing Chapter of this manual for download instructions.

- 2. Select Read from the Menu Bar.
- 3. Select either Detach.Batt info (Detachable Battery) or Inter.Batt info (Internal battery) depending on which battery you would like to read the cycle counts for.
- 4. Once selected the battery information will display in the Log window. The screen abbreviations are as follows:
 - Capacity Battery capacity displayed as a percentage.
 - V Battery Voltage displayed in millivolts.
 - I Battery Current displayed in miliamps
 - **T** Battery Temperature displayed in Celsius.
 - SH State of Health displayed as a percentage.
 - **CC** Battery Cycle Count displayed as a number between 0 and 500.
 - CF Condition Flag displayed as a 1 or 0.
 - ME Max Error displayed as a percentage.
 - S/N Battery Serial Number displayed as a number.
 - SM Ship Mode displays either Standby or Success.



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7.6 Internal Battery Conditioning Procedure

Respironics recommends replacement of the Internal Battery when the battery reaches 400 or more cycles. If the Internal Battery is not replaced and the Cycle Count is greater than 400 or the Max Error is greater than 25% then the following procedure must be performed to ensure it is still functioning properly.

- 1. Remove the Detachable Battery.
- 2. Set the device to the following therapy settings:
 - T mode
 - IPAP = 34 pressure units
 - EPAP = 4 pressure units
 - BPM = 10
 - Rise Time = 1
 - Inspiratory Time = 1 second
- 3. Connect AC, turn on Ventilator and charge battery to 100% capacity. Battery is fully charged when the Lightning bolt charge indicator is not displayed.
- 4. Disconnect AC and allow battery to discharge until Ventilator turns off.
- 5. Allow battery to rest for a minimum of 5.5 hours. Do not connect AC power and do not attempt to turn the Ventilator on.
- 6. Connect AC power and allow the Internal Battery to charge to 100%. Battery is fully charged when the Lightning bolt charge indicator is not displayed.
- 7. Allow battery to rest for minimum 2.5 hours. Do not disconnect AC power during this time.
- 8. Open the Trilogy Toolbox software and read the Internal Battery State of Health and MaxError using the Tool Box. Refer to Section 7.5 for more instructions.

NOTE

If the Trilogy Toolbox software is not downloaded to your computer, proceed to the Testing Chapter of this manual for download instructions.

9. The Internal Battery must have a State of Health ≥ 60% and MaxError =1%. If the Internal Battery does not meet the pass criteria it must be replaced.



7.7 CLEANING AND DISINFECTION PROCEDURES

7.7.1 CLEANING THE VENTILATOR

The ventilator's exterior surface and the exterior of the detachable battery pack (if using) should be cleaned before and after each patient use, and more often if needed.

WARNING

To avoid electrical shock, always unplug the power cord from the wall outlet or external battery before cleaning.

CAUTION

- Do not immerse the device in liquid or allow any liquid to enter the enclosure, inlet filter, or any opening. This may result in equipment damage.
- Do not use harsh detergents, abrasive cleaners, or brushes to clean the ventilator system. Use only the cleaning agents and methods described below.
- 1. Unplug the device and clean the front panel and exterior of the enclosure as needed using a clean cloth dampened with any of the following cleaning agents:
 - Water
 - Soapy water or a mild detergent
 - Hydrogen Peroxide (3%)
 - Isopropyl Alcohol (91%)
 - 10% bleach solution (10% bleach, 90% water)
- 2. Do not allow any liquid to drip into the ventilator case or detachable battery pack. After cleaning, use a soft, dry cloth to remove any residual cleaner. Use extra care when cleaning the display. Abrasive cleaners can scratch the display.
- 3. Allow the device to dry completely before plugging in the power cord.



7.7.2 CLEANING AND REPLACING THE AIR INLET FILTER

CAUTION

- Operating the device with a dirty filter may prevent the system from working properly and may damage the device.
- A dirty inlet filter may cause high operating temperatures that may affect device performance. Examine the inlet filter for integrity and cleanliness before each use. When the filter becomes dirty, it must be replaced to ensure normal device operation.

Under normal usage, you should clean the grey foam filter at least once every two weeks and replace it with a new filter every six months or sooner if needed.

- 1. If the device is operating, stop the airflow by pressing the On/Off button. Disconnect the device from the power source.
- 2. Remove the filter from the enclosure by gently squeezing the filter in the center and pulling it away from the device.
- 3. Examine the filter for cleanliness and integrity.
- 4. Wash the gray foam filter in warm water with a mild detergent. Rinse thoroughly to remove all detergent residue. Allow the filter to air dry completely before reinstalling it. If the foam filter is torn or damaged, replace it. (Only Respironics-supplied filters should be used as replacement filters.)

CAUTION

Never install a wet filter into the device. It is recommended that you clean the filter in the morning and alternate using the two foam filters provided with the system to ensure sufficient drying time for the cleaned filter.

Reinstall the filter.



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CHAPTER 8: REPAIR & REPLACEMENT

8.0 CHAPTER OVERVIEW

This chapter illustrates the replaceable components for the Trilogy Ventilators. Procedures for replacing the components are also provided in this chapter.

NOTE

Refer to the Testing Section for required testing after component replacement.

NOTE

Refer to the Repair Kits Section for proper repair kit identification.

WARNING

• To prevent electrical shock, disconnect the electrical supply before attempting to make any repairs to the Trilogy Ventilators.

CAUTION

- Electronic components used in this device are subject to damage from static electricity. Repairs made to this device must be performed only in an antistatic, ESD-protected environment.
- During all repair and replacement procedures, ensure that any connections that are broken during the process (fittings, tubing, and hoses) are reconnected securely.
- When using a leak detector, be careful that it does not come in contact with any electrical components.
- Using the Trilogy Toolbox Application, place the unit in "Ship Mode" prior to any disassembly of the unit. Otherwise, ensure that the Internal Battery connection is the first to be disconnected and last to be reconnected during re-assembly.



8.1 EXTERNAL COMPONENT REMOVAL/INSTALLATION

8.1.1 DETACHABLE BATTERY PACK REPLACEMENT



FIGURE 8-1: DETACHABLE BATTERY PACK REMOVAL/INSTALLATION

Removal

1. Lift the black handle and remove the Detachable Battery Pack from the ventilator using the pull tab.

Installation

1. With the pull tab on the Detachable Battery Pack oriented at the top, place the black tab into the slot located on the bottom of the Detachable Battery compartment and push the detachable battery pack into place.

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8.1.2 INLET FILTER REPLACEMENT

Removal

 Grasp the Inlet Filter and remove it from the Inlet Air Path Assembly (Trilogy 100 & Trilogy 200) or from the Oxygen Blending Module (Trilogy O₂ & Trilogy 202).

Installation

1. Place the Inlet Filter into the slot on the Inlet Air Path Assembly (Trilogy 100 & Trilogy 200) or the slot on the Oxygen Blending Module (Trilogy O₂ & Trilogy 202).

8.1.3 INLET AIR PATH ASSEMBLY REPLACEMENT (TRILOGY 100 & TRILOGY 200 ONLY)

Removal

1. Remove the four screws securing the Inlet Air Path Assembly to the ventilator.



FIGURE 8-2: AIR PATH ASSEMBLY SCREW LOCATIONS

2. Pull the Inlet Air Path Assembly up and away from the ventilator.

Installation

- 1. Place the Inlet Air Path Assembly into place on the ventilator.
- 2. Ensure the Air Path Foam is properly in place (Gray (foam) side down).
- 3. Secure the Inlet Air Path Assembly to the ventilator, tightening the four screws to 8 in-lbs.



8.1.4 OXYGEN BLENDING MODULE REPLACEMENT

Removal

- 1. Remove the four screws securing the Inlet Air Path Assembly to the ventilator.
- 2. Remove the Oxygen Blending Module tube from the port on the Rear Enclosure.
- 3. Remove the four pin connector from the slot on the Rear Enclosure.
- 4. Remove the Grommet from the hole in the Rear Enclosure.
- 5. Remove the Oxygen Blending Module from the Rear Enclosure.

Installation

- 1. Connect the four pin connector to the slot on the Rear Enclosure, and seat grommet into opening, verifying no wires are pinched.
- 2. Connect the Oxygen Blending Module tube to the port on the Rear Enclosure, verifying the tube is routed over top of the green tube.
- 3. Seat the Oxygen Blending Module onto the Rear Enclosure.
- 4. Secure the Oxygen Blending Module to the Rear Enclosure with four screws, tightening to 8 in-lbs.

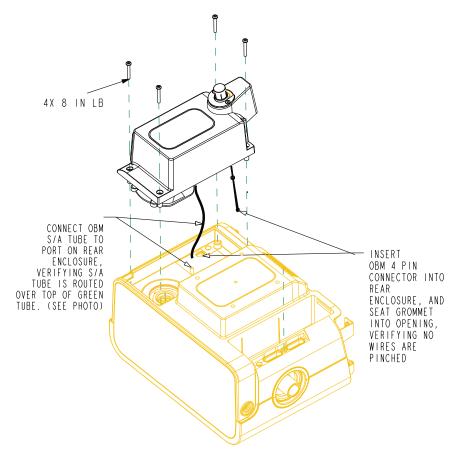


FIGURE 8-3: OXYGEN BLENDING MODULE REMOVAL/INSTALLATION

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8.1.5 AIR PATH FOAM REPLACEMENT



FIGURE 8-4: AIR PATH FOAM REMOVAL/INSTALLATION

Removal

- 1. Remove the Inlet Air Path Assembly or Oxygen Blending Module. Refer to the Inlet Air Path Assembly Section or Oxygen Module Assembly Section for more details.
- 2. Remove the Air Path Foam from the ventilator.

Installation

- 1. Place the Air Path Foam into the ventilator. Ensure that the grey (foam) side is facing down.
- 2. Install the Inlet Air Path Assembly or Oxygen Blending Module. Refer to the Inlet Air Path Assembly Section or Oxygen Module Assembly Section for more details.



8.1.6 ACTIVE/PASSIVE PORTING BLOCK REPLACEMENT

Removal

1. Remove the screw securing the Active/Passive Porting Block to the ventilator.



2. Lift the Active/Passive Porting Block straight up away from the ventilator.

Installation

- 1. Push the Active/Passive Porting Block into place.
- 2. Secure the Active/Passive Porting Block to the ventilator using one screw.

8.1.7 HANDLE REPLACEMENT

Removal

1. Remove the two 1/8" Hex screws securing the handle to the ventilator.



FIGURE 8-5: SCREW REMOVAL (REPEAT FOR OTHER SIDE)

- 2. Remove the Handle from the Ventilator.
- 3. Remove the two handle o-rings from the ventilator.

Installation

- 1. Insert the two handle o-rings in the ventilator.
- 2. Secure the handle to the ventilator by tightening the two 1/8" Hex screws to 8 in-lbs.

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8.2 FRONT/REAR/BOTTOM ENCLOSURE REMOVAL

CAUTION

Circuitry is powered if the Internal battery is connected.

- Remove the Detachable Battery Pack. Refer to the Detachable Battery Pack Replacement Section for more details.
- 2. Remove the Inlet Air Path Assembly or Oxygen Blending Module. Refer to the Inlet Air Path Assembly Section or Oxygen Blending Module Section for more details.
- 3. Remove the Air Path Foam. Refer to the Oxygen Blending Module Replacement Section for more details.
- 4. Remove the Active/Passive Porting Block. Refer to the Active/Passive Porting Block Replacement Section for more details.
- 5. Remove the Handle. Refer to the Handle Replacement Section for more details.
- 6. Lay the Trilogy on its top to expose the Bottom Enclosure.
- 7. Remove the four screws securing the Bottom Enclosure to the Trilogy.



FIGURE 8-6: BOTTOM ENCLOSURE SCREW LOCATIONS

- 8. With the LCD facing you lift up the Bottom Enclosure and remove the Alarm connections from Speaker 1 and Speaker 2 on the Front Panel Board.
- 9. Lay the ventilator LCD side down on a protected surface.
- 10. **IMPORTANT TO BE DISCONNECTED FIRST:** Remove the Internal Battery Cable from J2 on the Power Management Board.
- Remove the OBM wire harness from J7 on the Interface PCA (Trilogy O₂ and Trilogy 202 Only).
- 12. Remove the DC Battery Cable from J4 on the Power Management Board.
- 13. Remove the oxygen tube from the Exhalation Control Module.
- 14. Remove the Ethernet Cable from J19 on the System Board.
- 15. Remove the Interface Cable from J7 on the System Board.
- 16. Remove the AC Connector from J1 on the Power Supply Board.
- 17. Remove the Battery Fan Cable from J18 on the System Board.
- 18. Remove the Capacitor wires from J6 on the System Board.



- 19. Slide the Bottom Enclosure away from the Front and Rear Enclosures. Proceed to the Bottom Enclosure Assembly Component Removal/Installation section for instructions on how to remove/install the Bottom Enclosure Assembly components.
- 20. Remove the four screws securing the Front and Rear Enclosures together.



- 21. While holding the enclosures together place the Trilogy on its top so the opening is facing up.
- 22. Gently slide the Front and Rear Enclosures apart.
- 23. Remove the Sensor Cable from J4 on the Sensor Board.
- 24. Remove the Blower Cable from J5 on the System Board.
- 25. Remove the Blower Fan Cable from J17 on the System Board.
- 26. Remove the tape securing the Exhalation Control wires to the metal support.
- 27. Remove the Exhalation Control Module wires from J10 on the System Board.
- 28. Remove the Exhaust Fan wires from J1 on the System Board.
- 29. Remove the Detachable Battery Cable from J1 and J3 on the Power Management Board.
- 30. Cut the two tie wraps securing the Blower wires to the plastic support bracket.
- 31. Slide the Front and Rear Enclosures Apart.
- 32. Remove the Top Plate from the device.
- 33. The Front and Rear Enclosure are now disconnected. Proceed to the Front or Rear Enclosure Assembly Component Removal/Installation sections for instructions on how to remove/install the Assembly components. Also, Refer to the Oxygen Blending Module Components Section for instructions on how to remove/install the Oxygen Blending Module components.

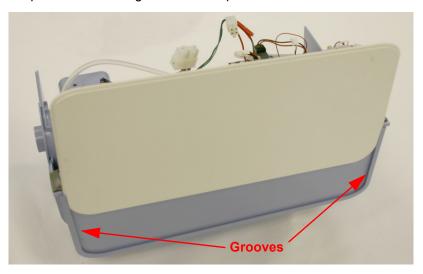
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8.2.1 TOP PLATE REMOVAL

Removal

- 1. Disconnect the Front, Rear, and Bottom Assemblies. Refer to the Front/Rear/Bottom Enclosure Removal Section for more details.
- 2. Slide the Top Plate out of the groves in the top of the Front or Rear Assemblies.



Installation

- 1. Slide the Top Plate into the groves in the top of the Front or Rear Assemblies.
- 2. Install the Front, Rear, and Bottom Assemblies. Refer to the Front/Rear/Bottom Enclosure Installation Section for more details.



8.3 BOTTOM ENCLOSURE ASSEMBLY COMPONENT REMOVAL/INSTALLATION

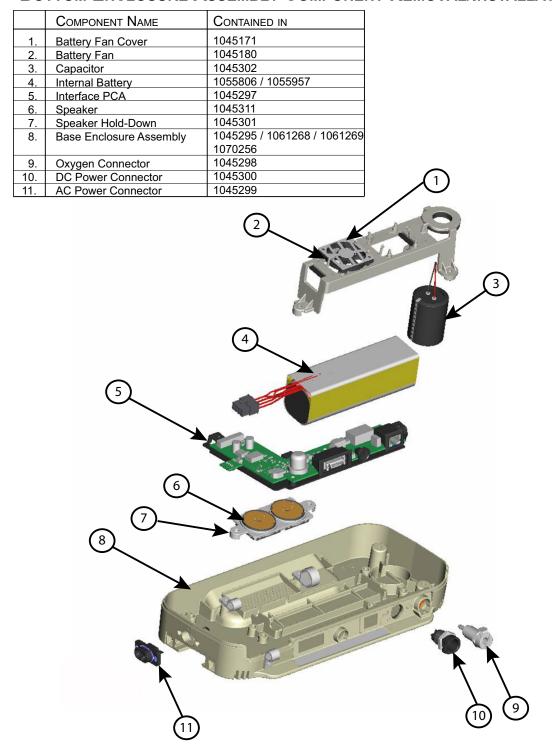


FIGURE 8-7: BOTTOM ENCLOSURE ASSEMBLY

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8.3.1 AC POWER CONNECTOR REPLACEMENT

Removal

Using a small phillips screwdriver and a 1/4" wrench, remove the two screws securing the AC Connector to the Bottom Enclosure. Discard the old screws and locking nuts if new screws and locking nuts are provided in your RP Kit.

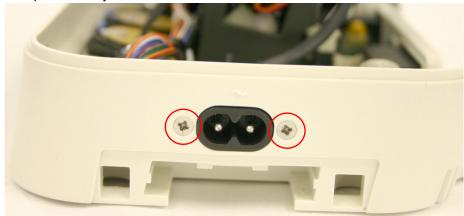


FIGURE 8-8: SCREW LOCATIONS

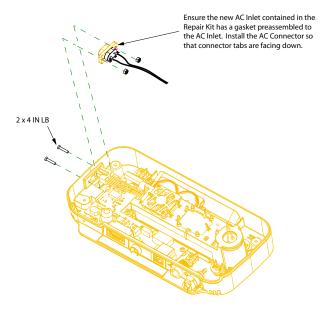
2. Remove the AC Power Connector from the Bottom Enclosure.

Installation

- 1. The new AC Inlet should have a new gasket attached to the plastic. Ensure that the Gasket is still in place.
- 2. Slide the AC Power Connector into the proper slot in the Bottom Enclosure. Slide the AC Power Connector into the proper slot in the Bottom Enclosure.



3. Using a 4 in-lbs torque wrench and a 1/4" wrench, secure the AC Connector to the Bottom Enclosure (new screws and locking nuts should be used if they are provided in RP Kit).



NOTE

If **ONLY** the AC Power Connector is replaced, you can use either the Performance Verification Tool located in the Maintenance Section of this manual or the Field Service Application located in the Testing Section of this manual as the Final Test Tool.

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8.3.2 DC POWER CONNECTOR REPLACEMENT

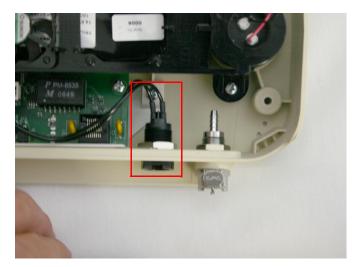


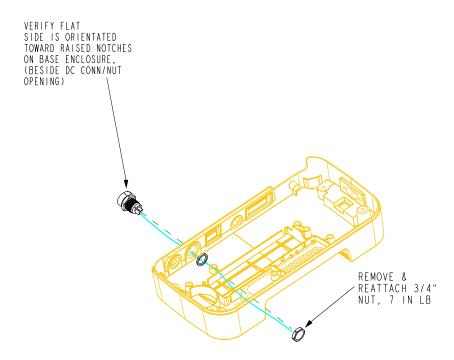
FIGURE 8-9: DC POWER CONNECTOR

Removal

- 1. Using a 3/4" wrench, remove the nut securing the DC Connector to the Bottom Enclosure.
- 2. Remove the DC Connector from the Bottom Enclosure.

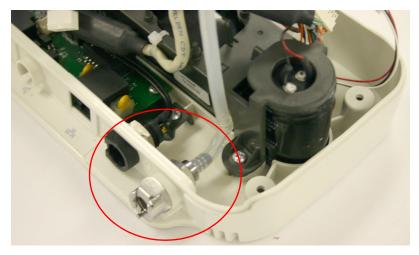
Installation

- 1. Place the DC Connector into its slot in the Bottom Enclosure.
- 2. Using a 3/4" wrench, connect the DC Connector to the Bottom Enclosure by tightening the nut to 7 in-lbs.





8.3.3 OXYGEN OUTLET REPLACEMENT

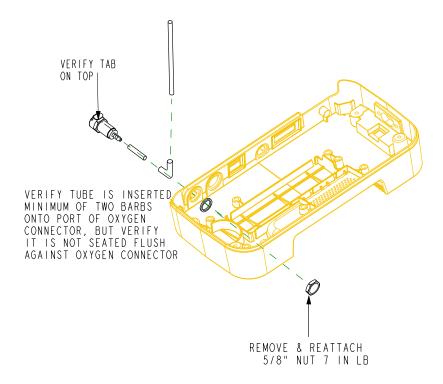


Removal

- 1. Using a 5/8" wrench, remove the nut securing the Oxygen Outlet to the Bottom Enclosure.
- 2. Remove Oxygen Outlet and tubing from the Bottom Enclosure.

Installation

- 1. Place the Oxygen Outlet into its slot in the Bottom Enclosure.
- 2. Connect the tubing to the Oxygen Outlet as shown below.
- 3. Using a 5/8" wrench, connect the Oxygen Outlet to the Bottom Enclosure by tightening the nut to 7 in-lbs.



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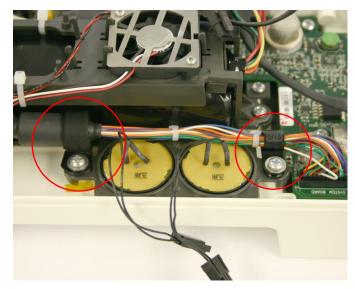
8.3.4 SPEAKER REPLACEMENT

Removal

- 1. Remove the two screws securing the Hold-Down and two cord clips to the Bottom Enclosure.
- 2. Remove the Speakers from the device.

Installation

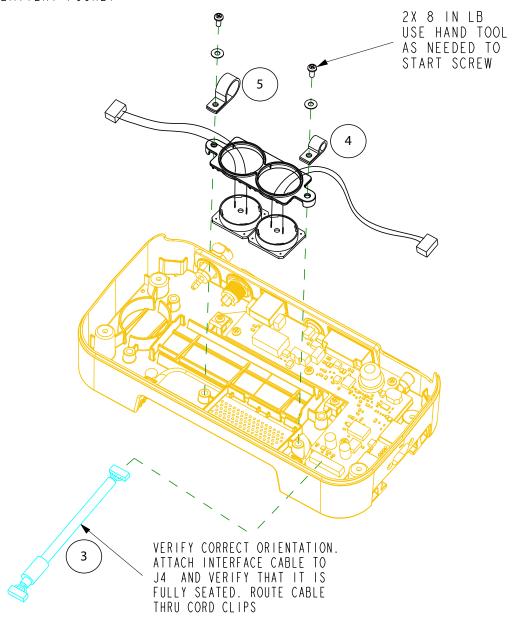
- 1. Place the Speaker in the Bottom Enclosure.
- 2. Place the Hold-Down into its position in the Bottom Enclosure.
- 3. With the Bottom Enclosure positioned with the AC Inlet on the right, place the cord clips in their proper position (the 5/16" cord clip should be positioned on the right and the larger cord clip on the left).



- 4. Ensure the Interface Cable is properly routed through the cord clips.
- 5. Secure the cord clips and hold-down to the Bottom Enclosure by tightening the two screws to 8 inlbs.



ROUTE ITEM 3 THRU ITEM 4 & 5 BEFORE TIGHTENING. FERRITE FITS INTO ITEM 5. CLIP LOOPS FACE TOWARD BATTERY POCKET



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8.3.5 INTERNAL BATTERY FAN REPLACEMENT

Removal

1. Unscrew the Battery Fan Cover from the Battery Fan.

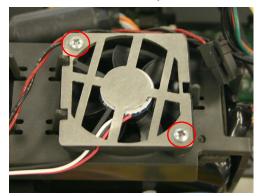


FIGURE 8-10: BATTERY FAN COVER

- 2. Cut the Tie warp securing the Fan wires to the Capacitor/Internal Battery Fan Retainer.
- 3. Release the locking tabs securing the Internal Battery Fan to the Capacitor/Internal Battery Retainer.

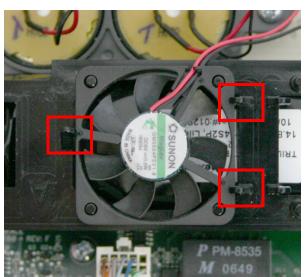


FIGURE 8-11: INTERNAL BATTERY FAN LOCKING TABS (BATTERY SHOWN WITHOUT BATTERY FAN COVER)

4. Remove the Internal Battery Fan from the Capacitor/Internal Battery Fan Retainer.

Installation

- 1. Align the holes in the Internal Battery Fan with the pins on the Retainer.
- 2. Push the Internal Battery Fan down until it locks in place.



3. Using a tie wrap secure the Fan wires to the Capacitor/Internal Battery Fan Retainer. The wires are to be secured to the front of the Capacitor/Battery Retainer with the wire tie in the orientation below. The wires must be loose and not pinched by the wire tie.

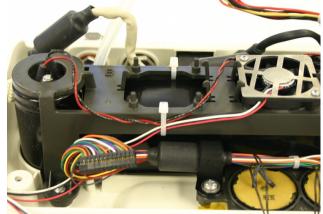
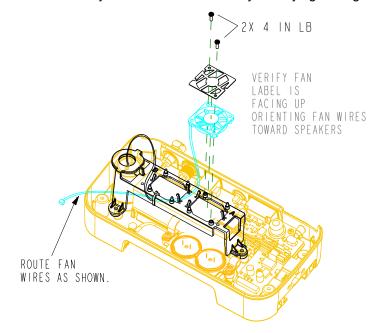


FIGURE 8-12: PROPER ROUTING OF FAN AND CAPACITOR WIRES

4. Secure the Battery Fan Cover to the Battery Fan by tightening it to 4 in-lbs.



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8.3.6 CAPACITOR/INTERNAL BATTERY REPLACEMENT

Removal

1. Remove the three screws securing the Capacitor/Internal Battery Retainer to the Bottom Enclosure.



FIGURE 8-13: CAPACITOR/INTERNAL BATTERY FAN RETAINER SCREWS

- 2. Cut the two tie wraps securing wires to the Capacitor/Internal Battery Retainer.
- 3. Remove the Capacitor and/or Internal Battery.

Installation

- 1. Place the Capacitor into its slot in the Bottom Enclosure.
- 2. Route the Internal Battery cable through the slot in the Internal Battery/Capacitor retainer.

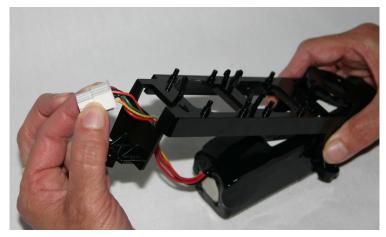
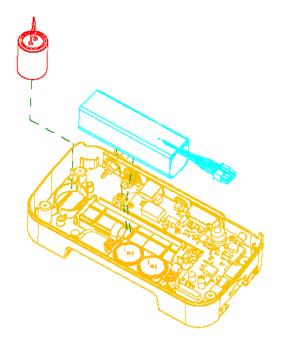
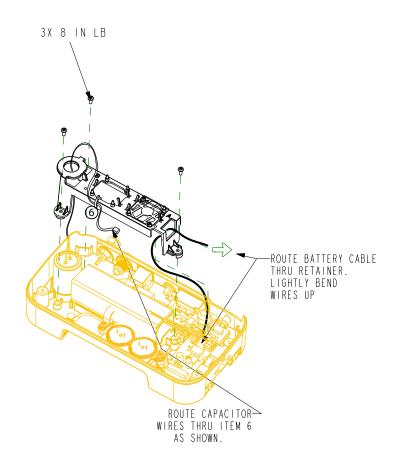


FIGURE 8-14: PROPER CABLE ROUTING

- 3. Secure the Capacitor and/or Internal Battery to the Bottom Enclosure using the Capacitor/Battery Retainer. Tighten the three screws to 8 in-lbs.
- 4. Using a tie wrap, secure the Capacitor wires to the Capacitor/Internal Battery Retainer.







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8.3.7 INTERFACE PCA REPLACEMENT

Removal

- 1. Remove the Internal Battery/Capacitor Retainer.
- 2. Remove the Interface to System Board Cable from J4 on the Interface PCA.
- 3. Remove the Ethernet to System Board Cable from J5 on the Interface PCA.



FIGURE 8-15: ETHERNET TO SYSTEM BOARD CABLE CONNECTION

4. Remove the four screws securing the Interface PCA to the Interface Board Retainer.

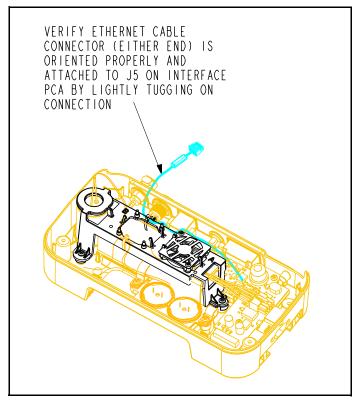


FIGURE 8-16: INTERFACE PCA SCREW LOCATIONS

Installation

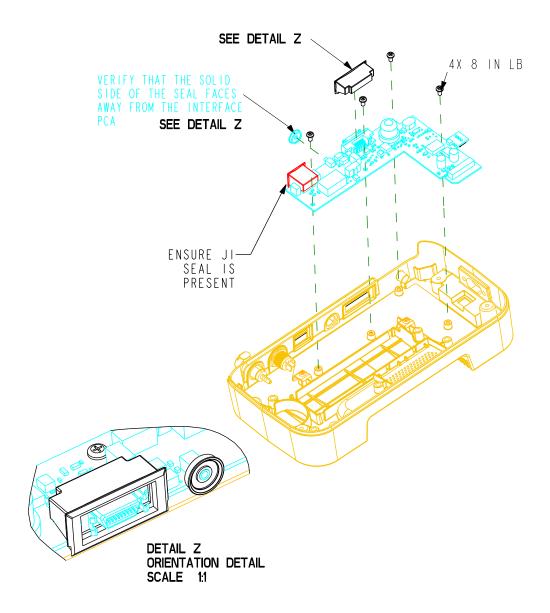
- 1. Secure the Interface PCA to the Bottom Enclosure by tightening the screws to 8 in-lbs.
- 2. Connect the Ethernet to System Board Cable to J5 on the Interface PCA.





- 3. Connect the Interface to System Board Cable from J4 on the Interface PCA.
- 4. Connect the Interface PCA to the Interface PCA Retainer.







8.3.8 INTERFACE PCA RETAINER REPLACEMENT

Removal

- 1. Remove to the Interface PCA. Refer to the Interface PCA Replacement section for more detailed instructions.
- 2. Remove the two screws securing the Interface PCA Retainer to the Bottom Enclosure.

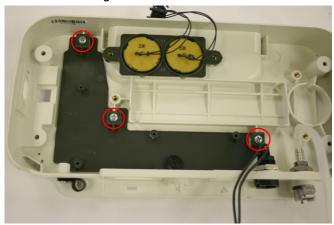
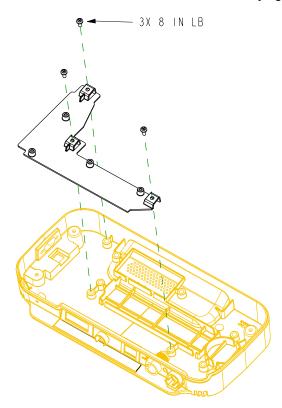


FIGURE 8-17: INTERFACE PCA RETAINER SCREW LOCATIONS

Installation

- 1. Place the Interface PCA Retainer into the Bottom Enclosure.
- 2. Secure the Interface Board Retainer to the Bottom Enclosure by tightening the screws to 8 in-lbs.



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8.3.9 BOTTOM ENCLOSURE REPLACEMENT

Removal

- 1. Remove the AC Power Connector. Refer to the Interface PCA Retainer Replacement Section for more details.
- Remove the DC Power Connector. Refer to the DC Power Connector Replacement Section for more details.
- 3. Remove the Oxygen Outlet. Refer to the Oxygen Outlet Replacement Section for more details.
- 4. Remove the Sonalert. Refer to the Speaker Replacement Section for more details.
- 5. Remove the Internal Battery Fan. Refer to the Internal Battery Fan Replacement Section for more details.
- 6. Remove the Capacitor and Internal Battery. Refer to the Capacitor/Internal Battery Replacement Section for more details.
- 7. Remove the Interface PCA. Refer to the Interface PCA Replacement Section for more details.
- Remove the Interface PCA retainer. Refer to the Interface PCA Retainer Replacement Section for more details.

Installation

- Install the Interface PCA retainer. Refer to the Interface PCA Retainer Replacement Section for more details.
- 2. Install the Interface PCA. Refer to the Interface PCA Replacement Section for more details.
- 3. Install the Capacitor and Internal Battery. Refer to the Capacitor/Internal Battery Replacement Section for more details.
- Install the Internal Battery Fan. Refer to the Internal Battery Fan Replacement Section for more details.
- 5. Install the Sonalert. Refer to the Speaker Replacement Section for more details.
- 6. Install the Oxygen Outlet. Refer to the Oxygen Outlet Replacement Section for more details.
- Install the DC Power Connector. Refer to the DC Power Connector Replacement Section for more details.
- Install the AC Power Connector. Refer to the Interface PCA Retainer Replacement Section for more details.
- 9. Print a generic Serial/Model Number label. The label must be in accordance with the following specifications: Label Size = 1-1/8" x 2" (maximum size), Font Size = 10 point (minimum)

IMPORTANT!

The new label MUST include the same Model Number, and Serial Number as those of which are on the original label.

- Affix the new label to the Bottom Enclosure in the same location as the original label. Affix the clear overlay (RI p/n 221013) over the printed label. Two clear overlays are provided with the Bottom Enclosure RP Kit.
- 11. Affix the Warning label in the same location as the original Warning Label.



8.4 FRONT ENCLOSURE ASSEMBLY COMPONENT REMOVAL/INSTALLATION

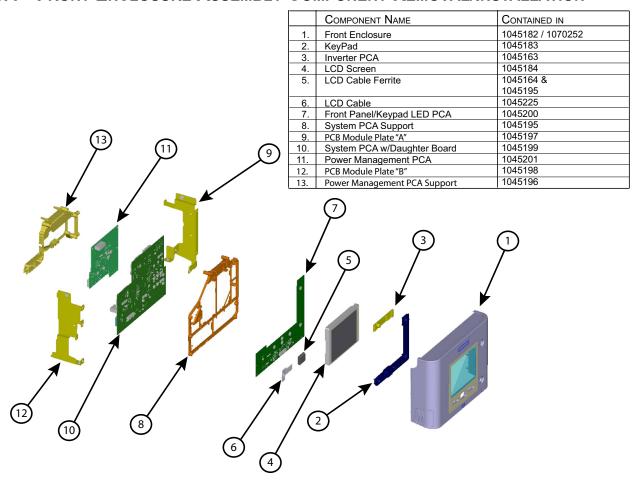


FIGURE 8-18: FRONT ENCLOSURE ASSEMBLY

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8.4.1 KEYPAD/FRONT ENCLOSURE REPLACEMENT

Removal

1. Remove the four screws securing the metal brackets to the front case. Three of the four screws are located between the metal bracket and the Front Enclosure.



FIGURE 8-19: METAL BRACKET SUPPORT SCREW LOCATIONS

- 2. Remove the assembly secured by the metal brackets out of the Front Enclosure to expose the Keypad.
- 3. Remove the Keypad.



FIGURE 8-20: KEYPAD REMOVAL



FIGURE 8-21: FRONT ENCLOSURE WITH KEYPAD REMOVED



Installation

- 1. Install the Keypad into the Front Enclosure.
- 2. Place the assembly secured with the metal brackets into the Front Enclosure.
- 3. Secure the Metal Brackets to the Front Enclosure by tightening the screw to 8 in-lbs.



8.4.2 FRONT PANEL PCA REPLACEMENT

Removal

1. Remove the 4 screws connecting the Front Panel PCA to the Power Management Board Support.

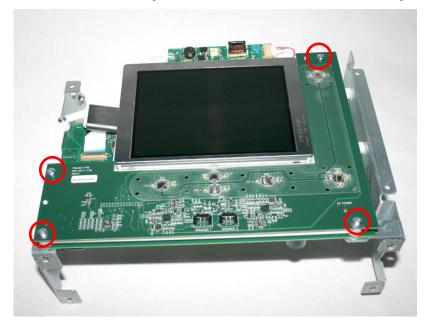


FIGURE 8-22: MOUNTING SCREW LOCATIONS

2. Remove the Front Panel PCA from the Assembly.

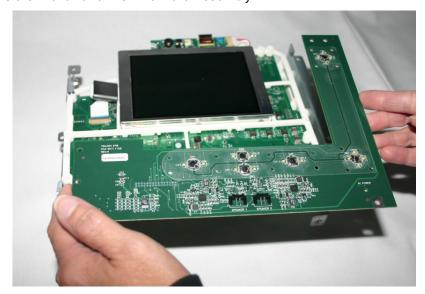


FIGURE 8-23: FRONT PANEL PCA REMOVAL

Installation

- 1. Place the Front Panel PCA on the Power Management Board Support.
- 2. Secure the Front Panel PCA to the Power Management Board Support by tightening the screws to 8 in-lbs. The two long screws go on the bottom and the two short screws go on the top.



8.4.3 PCA INVERTER REPLACEMENT

Removal

1. Remove the connection from location J20 on the Power Management Board.

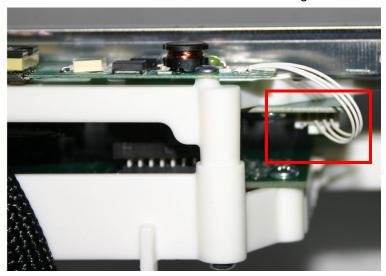


FIGURE 8-24: LOCATION J20 ON THE POWER MANAGEMENT PCA

2. Remove the LCD connection from location J2 on the PCA Inverter.

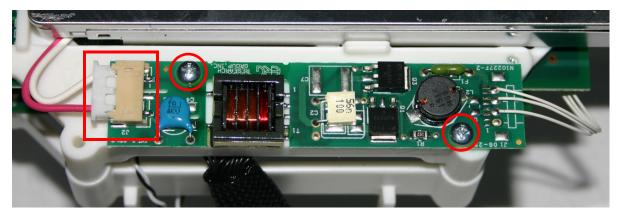


FIGURE 8-25: LOCATION J2 ON THE PCA INVERTER & SCREW LOCATIONS

3. Remove the two screws securing the PCA Inverter to the Power Management Board Support. Refer to Figure 8-25.

Installation

- 1. Secure the PCA Inverter to the Power Management Board Support by tightening the screws to 6 in-lbs.
- 2. Connect the LCD Cable to J2 on the PCA Inverter.
- 3. Connect the PCA Inverter to J20 on the Power Management Board.

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8.4.4 METAL SUPPORT BRACKET REPLACEMENT

Removal

- Remove the Front Panel PCA. Refer to the Front Panel PCA Replacement Section for more details.
- 2. Remove the PCA Inverter. Refer to the PCA Inverter Replacement Section for more details.
- 3. Remove the one remaining screw connecting the Power Management PCB Support to the Metal Bracket.



FIGURE 8-26: REMAINING SCREW

- 4. On a scratch proof surface, carefully turn the assembly over (LCD facing down).
- 5. Remove the 3 screws securing the PCB Supports to the side of the Metal Support Brackets.

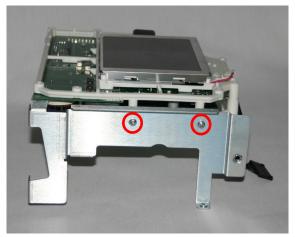




FIGURE 8-27: SCREWS SECURING THE METAL BRACKETS TO THE PCB SUPPORTS

Installation

- 1. Secure the PCB Supports to the side walls of the Metal Support Brackets by tightening the screws to 8 in-lbs.
- 2. Secure the Power Management PCB Support to the Metal Bracket.
- 3. Install the PCA Inverter. Refer to the PCA Inverter Replacement Section for more details.
- 4. Install the Front Panel PCA. Refer to the Front Panel PCA Replacement Section for more details.



8.4.5 POWER MANAGEMENT PCA REPLACEMENT

Removal

- 1. Remove the Keypad/Front Enclosure. Refer to the Keypad/Front Enclosure Replacement Section for more details.
- 2. Remove the Front Panel PCA. Refer to the Front Panel PCA Replacement Section for more details.
- 3. Remove the PCA Inverter. Refer to the PCA Inverter Replacement Section for more details.
- 4. Remove the Metal Support Brackets. Refer to the Metal Support Bracket Replacement Section for more details.
- 5. Remove the four screws securing the Power Management PCA support to the System PCA Support.



FIGURE 8-28: SCREW LOCATIONS

6. Depress the locking tab on the plastic standoff and firmly lift the Power Management PCA straight up and away from the System PCA.



FIGURE 8-29: LOCKING TAB

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7. Remove the four screws securing the Power Management PCA Support to the Power Management PCA.

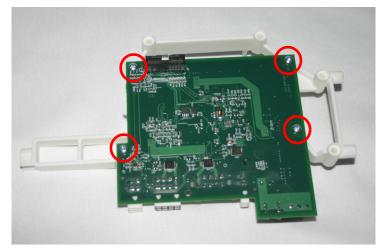


FIGURE 8-30: PCB SUPPORT SCREWS

Installation

- 1. Secure the Power Management PCA to the Power Management PCA Support by tightening the screws to 8 in-lbs.
- 2. Align the locking tab with the slot in the Power Management PCA. While firmly pressing down secure the Power Management PCA to the System PCA
- 3. Using the four screws, secure the Power Management PCA support to the System PCA support by tightening the screws to 12 in-lbs.
- 4. Install the Metal Support Brackets. Refer to the Metal Support Bracket Replacement Section for more details.
- 5. Install the PCA Inverter. Refer to the PCA Inverter Replacement Section for more details.
- 6. Install the Front Panel PCA. Refer to the Front Panel PCA Replacement Section for more details.
- 7. Install the Keypad/Front Enclosure. Refer to the Keypad/Front Enclosure Replacement Section for more details.



8.4.6 System PCA REPLACEMENT

IMPORTANT

After the System PCA has been replaced you must go to http://my.respironics.com to download the latest firmware to the Trilogy 100 Device. Follow the Operating Software Updates procedure located in the Testing Chapter of this Service Manual to complete the firmware installation. Once the latest firmware has been loaded to the device proceed to the Trilogy 100 Field Service Application for Final Testing of the device. If the firmware is not loaded the device will not operate or pass the Field Service Application Testing.

Removal

- Remove the Keypad/Front Enclosure. Refer to the Keypad/Front Enclosure Replacement Section for more details.
- 2. Remove the Front Panel PCA. Refer to Front Panel PCA Replacement Section for more details.
- 3. Remove the PCA Inverter.
- 4. Remove the Metal Support Brackets. Refer to Metal Support Bracket Replacement Section for more details.
- 5. Remove the Power Management PCA. Refer to the Power Management PCA Replacement Section for more details.
- 6. Lift up the brown tab and remove the LCD Cable from location J3 on the System PCA.

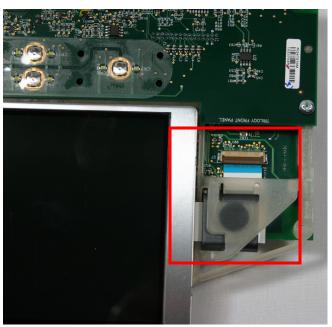


FIGURE 8-31: LCD CABLE AT LOCATION J3

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7. Remove the two screws securing the System PCA to the System PCA Support.



FIGURE 8-32: SCREW LOCATIONS

8. Remove the System PCA to Sensor PCA Cable from location J4 on the System PCA.

Installation

- 1. Connect the System PCA to Sensor PCA Cable to J4 on the System PCA.
- 2. Secure the Sensor PCA to the System PCA Support by tightening the screws to 8 in-lbs.
- 3. Connect the LCD Cable to J3 on the System PCA.
- 4. Install the Power Management PCA. Refer to the Power Management PCA Replacement Section for more details.
- 5. Install the Metal Support Brackets. Refer to the Metal Support Bracket Replacement Section for more details.
- 6. Install the PCA Inverter. Refer to the PCA Inverter Replacement Section for more details.
- 7. Install the Front Panel PCA. Refer to the Front Panel PCA Replacement Section for more details.
- Install the Keypad/Front Enclosure. Refer to the Keypad/Front Enclosure Replacement Section for more details.



8.4.7 LIQUID CRYSTAL DISPLAY (LCD) REPLACEMENT

Removal

- 1. Remove the Keypad/Front Enclosure. Refer to the Keypad/Front Enclosure Replacement Section for more details.
- 2. Remove Front Panel PCA. Refer to the Front Panel PCA Replacement Section for more details.
- 3. Remove the PCA Inverter. Refer to the PCA Inverter Replacement Section for more details.
- 4. Remove the Metal Support Brackets. Refer to the Metal Support Bracket Replacement Section for more details.
- 5. Remove the Power Management PCA. Refer to the Power Management PCA Replacement Section for more details.
- 6. Remove the System PCA. Refer to the System PCA Replacement Section for more details.
- 7. Lift up the brown locking tab, slide the cable through the ferrite, and remove LCD Cable from LCD.



FIGURE 8-33: LCD CABLE

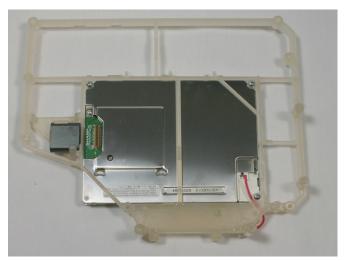


FIGURE 8-34: LCD CABLE REMOVED

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8. Remove the three screws securing the LCD to the System Board Support.

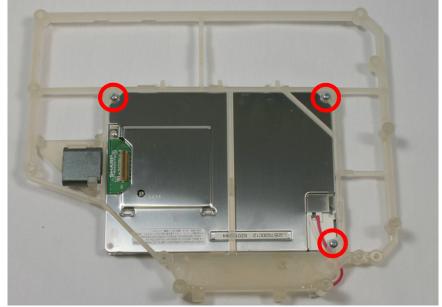


FIGURE 8-35: SCREW LOCATIONS

Installation

- 1. Slide the LCD Cable through the Ferrite and connect it to LCD.
- 2. Secure the LCD to the System Board Support by tightening the screws to 6 in-lbs.
- 3. Install the System PCA. Refer to the System PCA Replacement Section for more details.
- 4. Install the Power Management PCA. Refer to the Power Management PCA Replacement Section for more details.
- 5. Install the Metal Support Brackets. Refer to the Metal Support Bracket Replacement Section for more details.
- 6. Install the PCA Inverter. Refer to the PCA Inverter Replacement Section for more details.
- 7. Install the Front Panel PCA. Refer to the Front Panel PCA Replacement Section for more details.
- 8. Install the Keypad/Front Enclosure. Refer to the Keypad/Front Enclosure Replacement Section for more details.



8.5 REAR ENCLOSURE ASSEMBLY COMPONENT REMOVAL/INSTALLATION

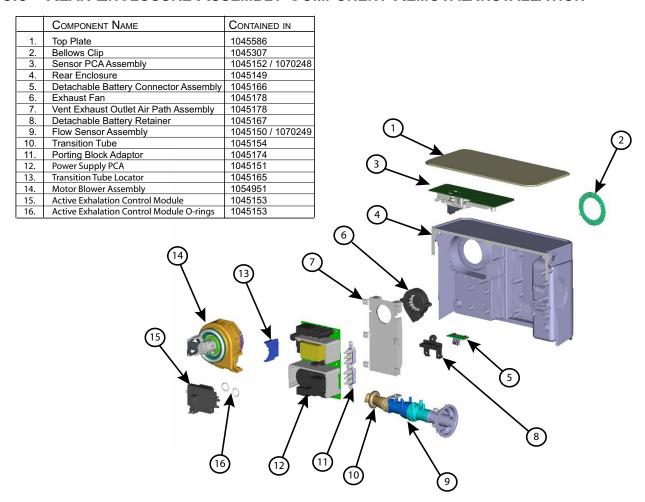


FIGURE 8-36: REAR ENCLOSURE ASSEMBLY

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8.5.1 DETACHABLE BATTERY CONNECTOR ASSEMBLY REPLACEMENT

Removal

- 1. Remove the connection from J2 located on the Trilogy Power Supply Board.
- 2. Remove the three screws connecting the Detachable Battery Connector Assembly to the Rear Enclosure.

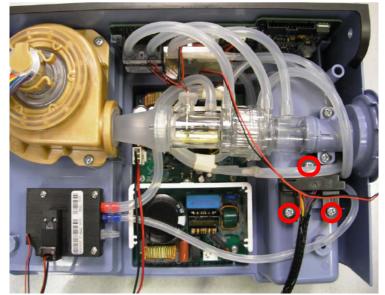


FIGURE 8-37: DETACHABLE BATTERY CONNECTOR SCREW LOCATIONS

3. Separate the Retainer from the Detachable Battery Assembly.

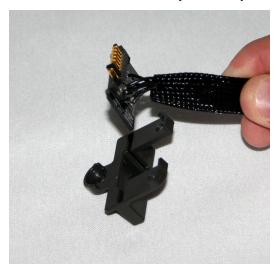


FIGURE 8-38: SEPARATING THE DETACHABLE BATTERY ASSEMBLY FROM THE RETAINER

Installation

- 1. Slide the Detachable Battery Connector Assembly into the Retainer.
- 2. Secure the Detachable Battery Connector Assembly to the Rear Enclosure by tightening the screws to 8 in-lbs.
- 3. Connect the two-prong connector to location J2 on the Trilogy Power Supply Board.



8.5.2 STIRRING FAN/STIRRING FAN RETAINER REPLACEMENT

Removal

- 1. Lift the center tab straight up to release the adhesive on the back of the tab from the Motor Blower Assembly.
- 2. Pull the Stirring Fan Retainer away from the Motor Blower Assembly.



FIGURE 8-39: STIRRING FAN RETAINER REMOVAL

3. Push the tabs on Stirring Fan Retainer out to remove the Stirring Fan from the Retainer.

Installation

- 1. Remove the adhesive backing from the Stirring Fan Retainer.
- 2. Clip the Stirring Fan Retainer onto the Motor Blower Assembly.
- 3. Place the Stirring Fan into the Stirring Fan Retainer.

NOTE

Verify that the Fan label is toward the Motor.

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8.5.3 ACTIVE EXHALATION MODULE REPLACEMENT

Removal

1. Remove the tubing from the Exhalation Module ports.

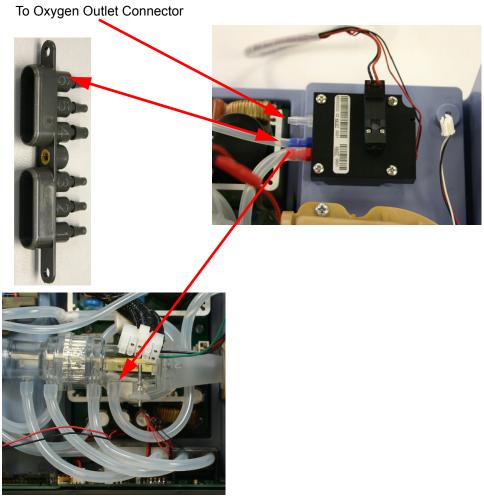


FIGURE 8-40: TUBING CONNECTIONS AND SCREW LOCATIONS

2. Remove the four screws securing the Active Exhalation Module to the Rear Enclosure.



Installation

1. Ensure the two O-rings are properly in place on the Active Exhalation Module.

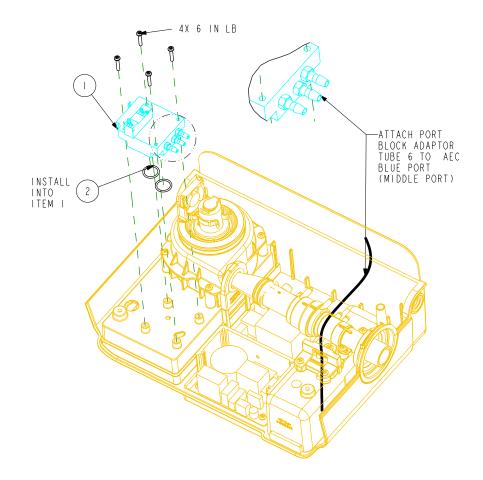


FIGURE 8-41: REMOVAL/INSTALLATION OF ACTIVE EXHALATION MODULE

- 2. Connect the Active Exhalation Module to the Rear Enclosure by tightening the four screws to 6 inlbs.
- 3. Connect the tubing to the Active Exhalation Module ports.



8.5.4 MOTOR BLOWER ASSEMBLY REPLACEMENT

Removal

1. Remove the three screws securing the Isolation Assembly to the Rear Enclosure.

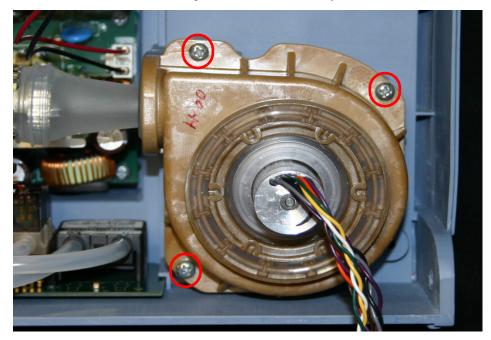


FIGURE 8-42: SCREW LOCATION

- 2. Slightly lift the entire Motor Blower Assembly up and out of the Rear Enclosure.
- 3. Disconnect the Transition Tube from the Flow Element Assembly.

Installation

- 1. Connect the Transition Tube to the Flow Element Assembly. Refer to Figure 8-43 for more detailed instructions.
- 2. Place the Assembly into the Rear Enclosure.
- 3. Secure the Assembly to the Rear Enclosure by tightening the screws to 8 in-bs.
- 4. Once the Motor Blower Assembly is secured, turn the enclosure over and wrap the Bellow around the Blower until it seals.



8.5.5 TRANSITION TUBE REPLACEMENT

Removal

- 1. Remove the Motor Blower Assembly. Refer to the Motor Blower Assembly Replacement Section for more details.
- 2. Remove the Transition Tube from the Flow Sensor Assembly.



FIGURE 8-43: DISCONNECTING/INSTALLING THE TRANSITION TUBE FROM FLOW SENSOR ASSEMBLY

Installation

- 1. Connect the small end of the Transition Tube to the Motor Blower Assembly. Verify proper alignment of the Transition Tube.
- 2. Connect the large end of the Transition Tube around the end of the Flow Sensor Assembly. Verify proper alignment of the Transition Tube.
- 3. Seat the Transition Tube disk into the Locator.
- 4. Install the Motor Blower Assembly. Refer to the Motor Blower Assembly Replacement Section for more details.

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8.5.6 TRILOGY 100 FLOW SENSOR ASSEMBLY REPLACEMENT

Removal

- 1. Remove the all of the tubing, including the Transition Tube, from the Flow Sensor Assembly.
- 2. Remove the two screws securing the Flow Sensor Assembly to the Rear Enclosure.

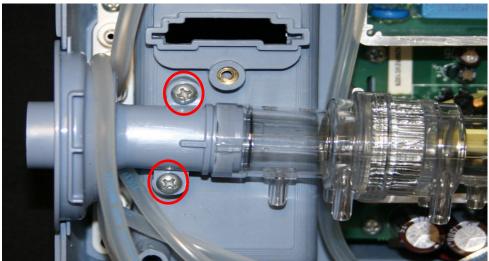


FIGURE 8-44: SCREW LOCATION

- 3. Remove the Outlet Flow Path Thermistor and Temperature Sensor Caps from the Flow Sensor Assembly.
- 4. Pull the Flow Sensor Assembly from the Rear Enclosure.



Installation

1. Ensure the two porting block caps are installed on the Flow Sensor Assembly.

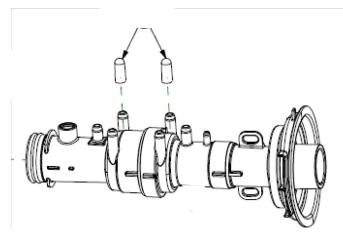


FIGURE 8-45: PORTING BLOCK CAP INSTALLATION

2. Place the Flow Sensor Assembly into the correct location in the Rear Enclosure.

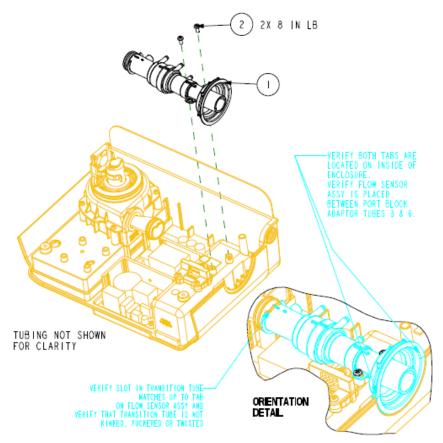


FIGURE 8-46: FLOW SENSOR ASSEMBLY INSTALLATION

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3. Place the Temperature Sensor Cap and Outlet Flow Path Thermistor into the proper location in the Flow Sensor Assembly.

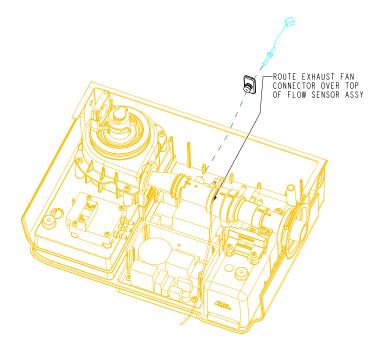


FIGURE 8-47: SENSOR CAP INSTALLATION

4. Secure the Flow Sensor Assembly to the Rear Enclosure by tightening the screws to 8 in-lbs.



5. Install all of the tubing back to the proper location. Refer to the drawings below.

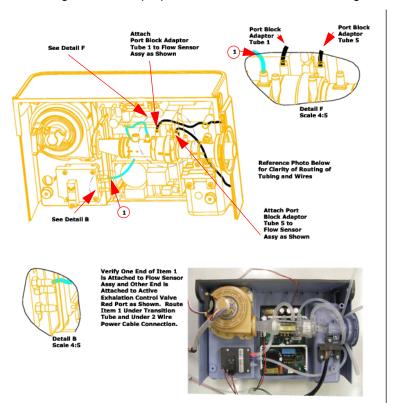


FIGURE 8-48: PROPER TUBING CONNECTIONS 1

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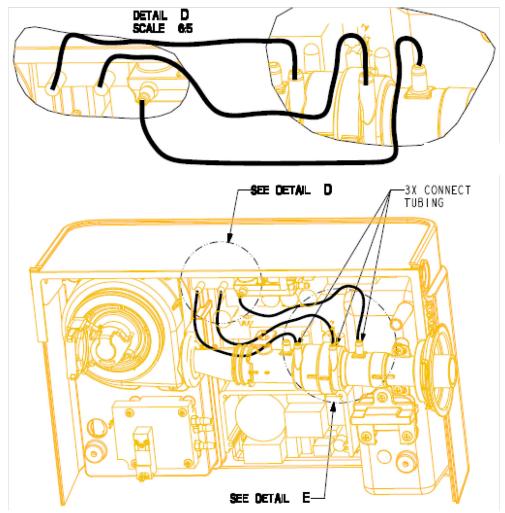


FIGURE 8-49: PROPER TUBING CONNECTIONS 2



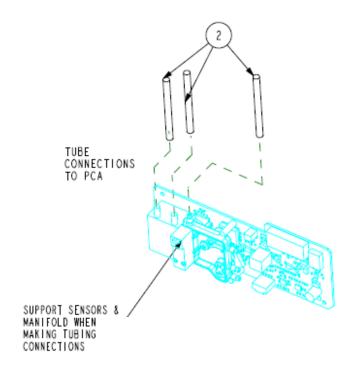


FIGURE 8-50: PROPER TUBING CONNECTIONS 3



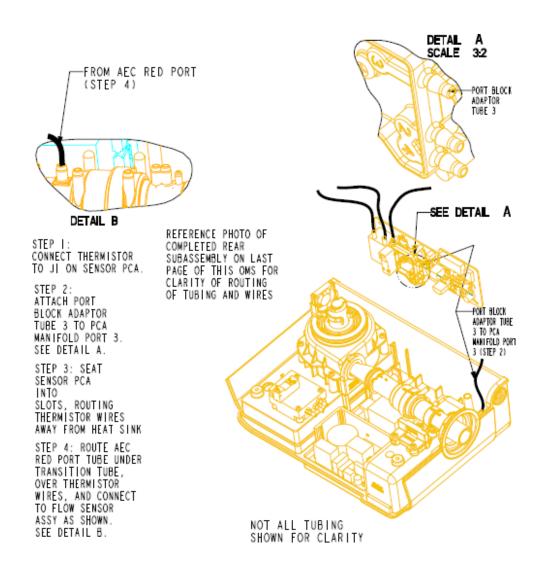


FIGURE 8-51: PROPER TUBING CONNECTIONS 4



8.5.7 TRILOGY 200, TRILOGY O2, & TRILOGY 202 FLOW SENSOR ASSEMBLY REPLACEMENT

Removal

- 1. Remove the all of the tubing, including the Transition Tube, from the Flow Sensor Assembly.
- 2. Remove the two screws securing the Flow Sensor Assembly to the Rear Enclosure.

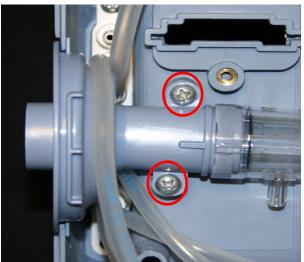


FIGURE 8-52: SCREW LOCATION

- 3. Remove the Outlet Flow Path Thermistor and Temperature Sensor Cap from the Flow Sensor Assembly.
- 4. Pull the Flow Sensor Assembly from the Rear Enclosure.

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Installation

1. Ensure the two porting block caps are installed on the Flow Sensor Assembly.

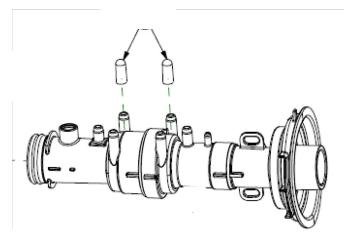


FIGURE 8-53; PORTING BLOCK CAP INSTALLATION

2. Place the Flow Sensor Assembly into the correct location in the Rear Enclosure.

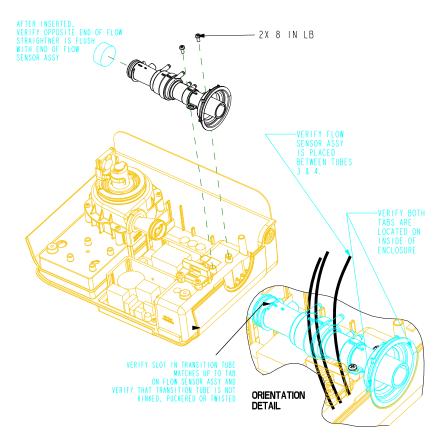


FIGURE 8-54: FLOW SENSOR INSTALLATION



3. Place the Temperature Sensor Cap and Outlet Flow Path Thermistor into the proper location in the Flow Sensor Assembly.

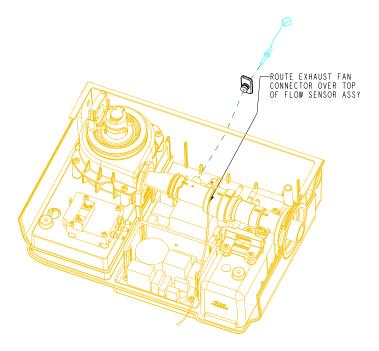


FIGURE 8-55: TEMPERATURE SENSOR CAP INSTALLATION

4. Secure the Flow Sensor Assembly to the Rear Enclosure by tightening the screws to 8 in-lbs.

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5. Install all of the tubing back to the proper location. Refer to the drawings below.

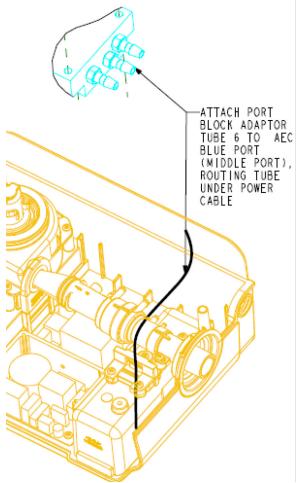


FIGURE 8-56: PROPER TUBING CONNECTIONS 1



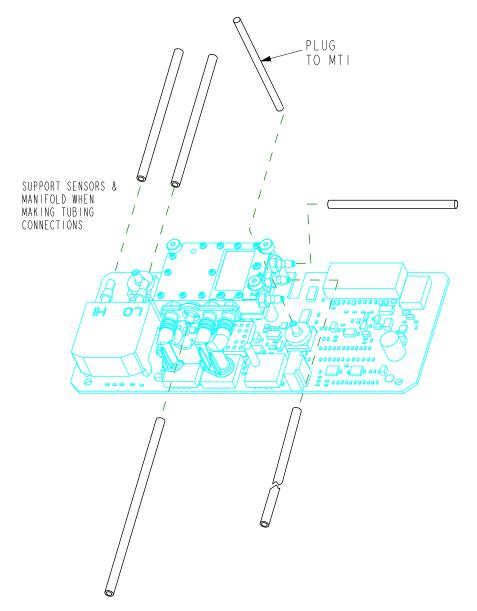


FIGURE 8-57: PROPER TUBING CONNECTIONS 2

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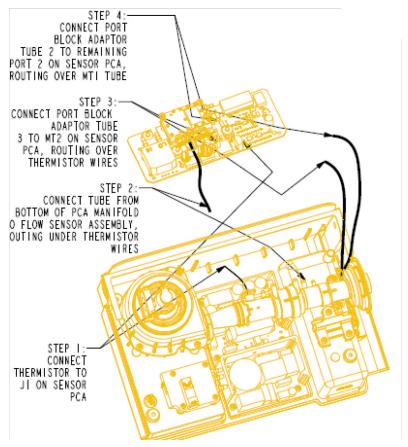


FIGURE 8-58: PROPER TUBING CONNECTIONS 3



8.5.8 PORTING BLOCK ADAPTOR REPLACEMENT

Removal

- 1. Remove the Flow Sensor Assembly. Refer to the Trilogy 100 Flow Sensor Assembly Section or the Trilogy 200, Trilogy O₂, & Trilogy 202 Section for more information for more details.
- 2. Remove all of the tubing connected to the Porting Block Adaptor.
- 3. Remove the two screws connecting the Porting Block Adaptor to the Rear Enclosure.

Installation

1. For Trilogy 100 device ensure the porting block caps are installed as shown.

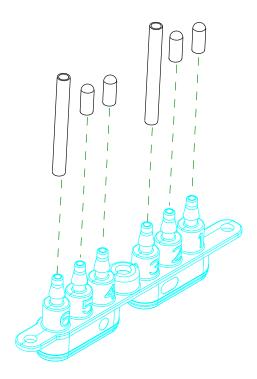


FIGURE 8-59: TRILOGY 100 TUBING & CAP PLACEMENT

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2. For Trilogy 200, Trilogy O₂, & Trilogy 202 devices ensure the porting block caps are installed as shown.

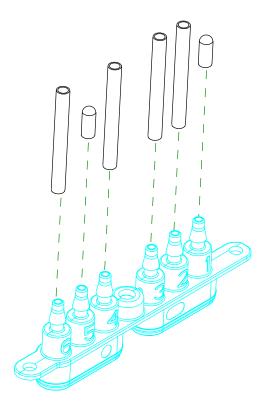


FIGURE 8-60: TRILOGY 200, TRILOGY O2, & TRILOGY 202 TUBING & CAP PLACEMENT



3. Connect the Porting Block Adaptor to the Rear Enclosure by tightening the screws to 8 in-lbs.

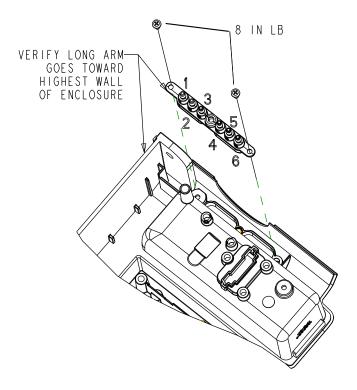


FIGURE 8-61: PORTING BLOCK INSTALLATION

NOTE

Verify that the long arm of the Porting Block Adaptor goes towards the highest wall of the enclosure.

- 4. Connect the correct tubing to the proper location on the Porting Block Adaptor.
- 5. Install the Flow Sensor Assembly. Refer to the Trilogy 100 Flow Sensor Assembly Section or the Trilogy 200, Trilogy O₂, & Trilogy 202 Section for more information for more details.

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8.5.9 TRILOGY 100 SENSOR PCA REPLACEMENT

Removal

1. Slide the Sensor PCA out of the Rear Enclosure.

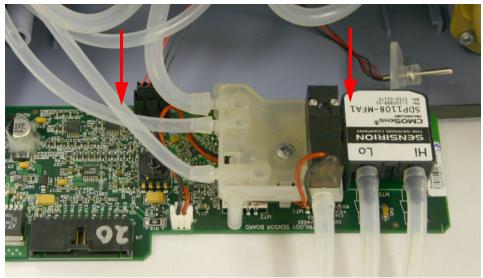


FIGURE 8-62: SENSOR PCA REMOVAL

2. Remove any tubing connected to the Sensor PCA.

Installation

1. Slide the Sensor PCA into the grooves in the Rear Enclosure.

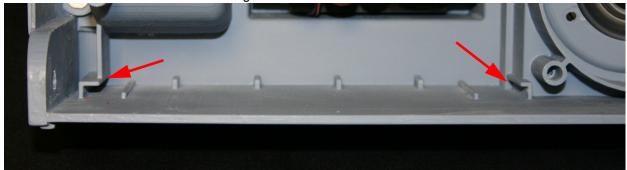


FIGURE 8-63: GROOVE LOCATION

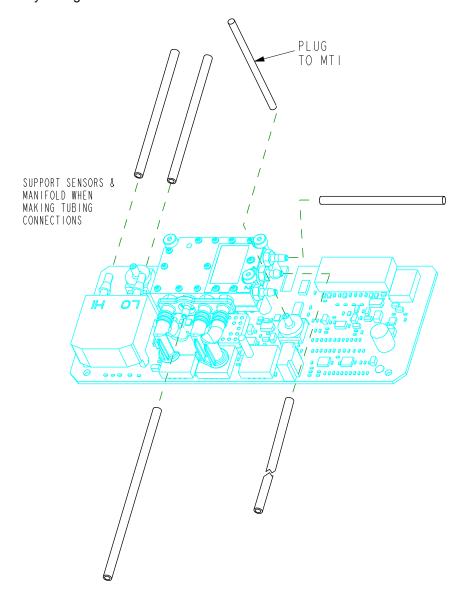
2. Connect the proper tubing to the Sensor PCA. Refer to Figure 8-50.



8.5.10 TRILOGY 200, TRILOGY O_2 , & TRILOGY 202 SENSOR PCA REPLACEMENT

Removal

- 1. Slide the Sensor PCA out of the Rear Enclosure.
- 2. Remove any tubing connected to the Sensor PCA.



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Installation

- 1. Connect thermistor to J1 on Sensor PCA.
- Connect tube from bottom of PCA Manifold to Flow Sensor Assembly, routing under the thermistor wires
- 3. Connect the Porting Block Adaptor tube 3 to MT2 on Sensor PCA, routing over thermistor wires.
- 4. Connect Port Block Adaptor Tube 2 to remaining port on Sensor PCA, routing over MT1 Tube.

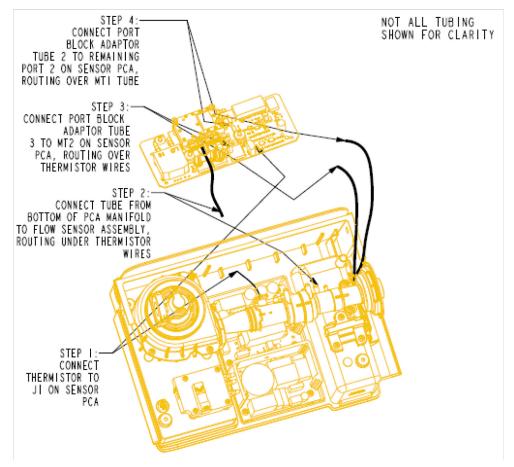


FIGURE 8-64: TUBING AND ELECTRICAL CONNECTIONS



Follow the steps in the Figure below to insert the PCA in the enclosure and make the final connections.

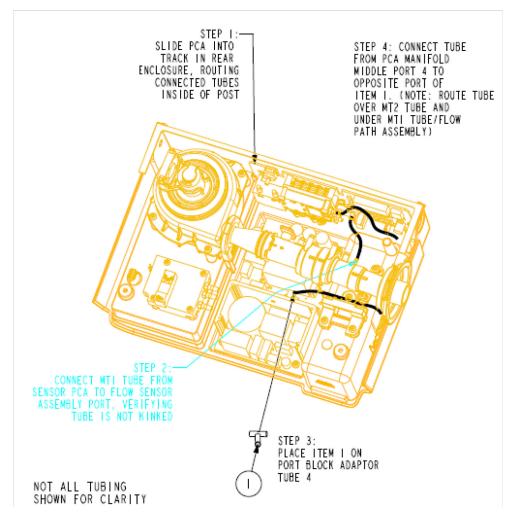


FIGURE 8-65: PCA PLACEMENT AND FINAL CONNECTIONS8



6. Ensure items 1, 2, 3, and 4 are assembled properly.

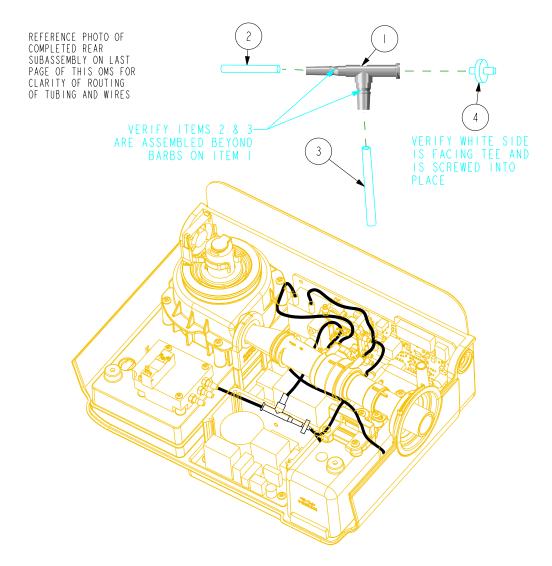


FIGURE 8-66: TEE TUBING CONNECTIONS

- 7. Ensure part 2 is attached to the Active Exhalation Control Valve red port.
- 8. Ensure item 3 is connected from the Filter/Tee to the Flow Sensor Assembly, routing the tube under the Flow Path Assembly and over the thermistor.
- 9. Connect the Tube from the Sensor PCA Manifold top port to filter, routing over MT2 tube and under MT1 tube and Flow Path Assembly.
- 10. Ensure the coiled pice of tubing has a plug attached to the end.
- 11. Ensure the coiled tube is attached to the Tee and nested under the Flow Sensor Assembly, with tubing routed under the filter/Tee Assembly.
- 12. Connect Sensirion Hi and Lo tubes to Outlet Flow Path.



8.5.11 POWER SUPPLY PCA REPLACEMENT

Removal

- Remove the Motor Blower Assembly. Refer to the Motor Blower Assembly Replacement Section for more details.
- 2. Remove the Transition Tube. Refer to the Transition Tube Replacement Section for more details.
- 3. Remove the Flow Sensor Assembly. Refer to the Trilogy 100 Flow Sensor Assembly Section or the Trilogy 200, Trilogy O_2 , & Trilogy 202 Section for more information for more details.
- Remove the Sensor PCA. Refer to the Trilogy 100 Sensor PCA Section or the Trilogy 200 & O₂ Sensor PCA Section for more details.
- 5. Remove the Power Supply PCA to Power Management Board Cable from location J2 on the Power Supply PCA.
- 6. Remove the six screws securing the Power Supply PCA to the Rear Enclosure.



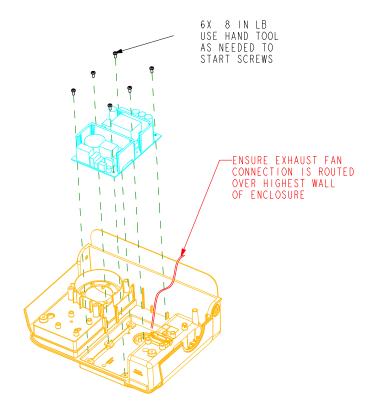
FIGURE 8-67: SCREW LOCATION

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Installation

1. Connect the Power Supply to the Rear Enclosure by tightening the screws to 8 in-lbs.



- 2. Connect the Power Supply PCA to Power Management Board Cable to location J2 on the Power Supple PCA.
- 3. Install the Sensor PCA. Refer to the Trilogy 100 Sensor PCA Section or the Trilogy 200 & O₂ Sensor PCA Section for more details.
- 4. Install the Flow Sensor Assembly. Refer to the Trilogy 100 Flow Sensor Assembly Section or the Trilogy 200, Trilogy O₂, & Trilogy 202 Section for more information for more details.
- 5. Install the Transition Tube. Refer to the Transition Tube Replacement Section for more details.
- 6. Install the Motor Blower Assembly. Refer to the Motor Blower Assembly Replacement Section for more details.



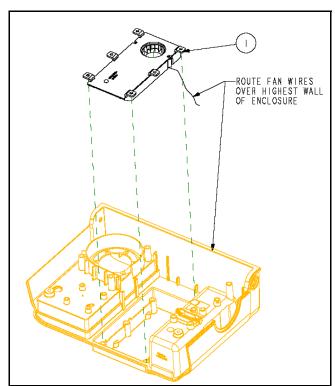
8.5.12 EXHAUST FAN ASSEMBLY REPLACEMENT

Removal

- 1. Remove the Motor Blower Assembly. Refer to the Motor Blower Assembly Replacement Section for more details.
- Remove the Transition Tube. Refer to the Transition Tube Replacement Section for more details.
- 3. Remove the Flow Sensor Assembly. Refer to the Trilogy 100 Flow Sensor Assembly Section or the Trilogy 200 & Trilogy O₂, & Trilogy 202 Section for more information for more details.
- Remove the Sensor PCA. Refer to the Trilogy 100 Sensor PCA Section or the Trilogy 200 & O₂ Sensor PCA Section for more details.
- Remove the Power Supply PCA. Refer to the Power Supply PCA Replacement Section for more details.
- 6. Lift the Exhaust Fan Assembly up and out of the Rear Enclosure.

Installation

1. Place the Exhaust Fan Assembly in the Rear Enclosure. Route the Fan wires over the highest wall of the enclosure.



- 2. Place the Power Supply PCA over top of the Exhaust Fan Assembly and secure it to the Rear Enclosure by tightening the screws to 8 in-lbs.
- 3. Connect the Power Supply PCA to Power Management Board Cable to location J2 on the Power Supple PCA.
- 4. Install the Sensor PCA. Refer to the Trilogy 100 Sensor PCA Section or the Trilogy 200 & O₂ Sensor PCA Section for more details.
- 5. Install the Flow Sensor Assembly. Refer to the Trilogy 100 Flow Sensor Assembly Section or the Trilogy 200, Trilogy O₂, Trilogy 202 Section for more information for more details.

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- 6. Install the Transition Tube. Refer to the Transition Tube Replacement Section for more details.
- 7. Install the Motor Blower Assembly. Refer to the Motor Blower Assembly Replacement Section for more details.

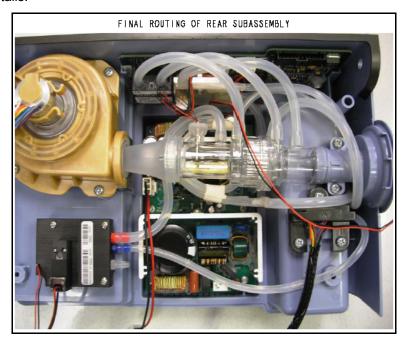


FIGURE 8-68: PROPER ROUTING OF TRILOGY 100 TUBING AND WIRING

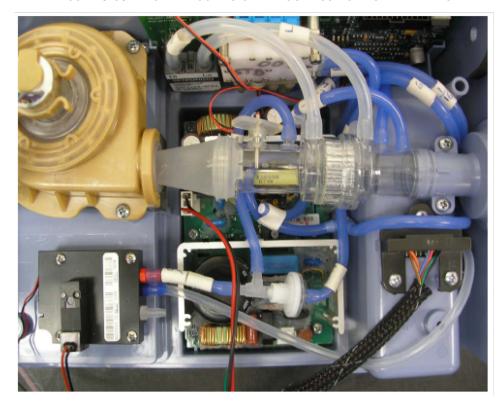


FIGURE 8-69: PROPER ROUTING OF TRILOGY 200, TRILOGY O2, TRILOGY 202 TUBING AND WIRING



8.6 FRONT/REAR/BOTTOM ENCLOSURE INSTALLATION

- IMPORTANT TO BE CONNECTED LAST: Connect the Internal Battery Cable to J2 on the Power Management Board
- 2. Place the Top Plate in the groves on the Front and Rear Enclosure.
- 3. Move the enclosure close enough to connect the wiring.
- 4. Using two tie wraps, secure the Ferrite on the Blower wires to the plastic support bracket.
- 5. Connect the Detachable Battery cable to J1 and J3 on the Power Management Board.
- 6. Connect the Exhaust Fan to J1 on the System Board.
- 7. Connect the Exhalation Control Module to J10 on the System Board.
- 8. Secure the Exhalation Control Module Wires to the Metal Support using cloth tape.
- 9. Connect the Blower Fan to J17 on the System Board.
- 10. Connect the Blower to J5 on the System Board.
- 11. Connect the Sensor Cable to J4 on the Sensor Board.
- 12. Ensuring not to pinch tubing or wiring, slide the Front and Rear Enclosures completely together.
- 13. Secure the Front and Rear Enclosures together by tightening the screws to 8 in-lbs.
- 14. The Front and Rear Enclosure are now in place. It is now time to connect the Bottom Enclosure.
- 15. Connect the Capacitor to J6 on the System Board.
- 16. Connect the Battery Fan Cable to J18 on the System Board.
- 17. Connect the AC Connector to J1 on the Power Supply Board.
- 18. Connect the Power Supply to location J1 on the Power Management Board.
- 19. Connect the Interface Cable to J7 on the System Board.
- 20. Connect the Ethernet Cable to J19 on the System Board.
- 21. Connect the Oxygen Tube to the Exhalation Control Module.
- 22. Connect the DC Battery Cable to J4 on the Power Management Board.
- 23. Connect the OBM wire harness to J7 on the Interface PCA (Trilogy O₂ and Trilogy 202 Only).
- 24. **IMPORTANT TO BE CONNECTED LAST:** Connect the Internal Battery Cable to J2 on the Power Management Board.

CAUTION

Once the Internal Battery Cable is connected, power may be active.

- 25. Ensure all wiring connections are still connected and the Oxygen Tube is still connected.
- 26. Fold the Bottom Enclosure over to seal the enclosure together. Ensure that the wiring and tubing is clear of the seam and not kinked.
- 27. Lift up the device and turn the device to have the LCD face you. Gently lift up one end of the Bottom Enclosure and connect the Alarms to Speaker 1 and Speaker 2 on the front Panel Board. Ensure that the wiring and tubing is clear of the seam and not kinked.
- 28. Secure the Bottom Enclosure to the Front and Rear Enclosures by tightening the screws to 8 inlbs.
- 29. Connect the Active/Passive Port. Refer to the Active/Passive Porting Block Replacement Section for more details.
- 30. Insert the Air Path Foam. Refer to the Air Path Foam Replacement Section for more details.



31. Connect the Inlet Air Path Assembly or Oxygen Blending Module Assembly. Refer to the Inlet Air Path Assembly Section or the Oxygen Blending Module Assembly procedure for more details.

8.7 OXYGEN BLENDING MODULE COMPONENT REPAIR & REPLACEMENT PROCEDURES

8.7.1 OXYGEN BLENDER FILTER/WHISPER CAP REPLACEMENT

Removal

- 1. Pull the Whisper Cap from the Air Inlet Duct.
- 2. Remove the Filter from the Air Inlet Duct.

Installation

- 1. Place the Filter into the Air Inlet Duct.
- 2. Place the Filter Cap over the Filter and ensure it is securely in place (you will hear a snap).

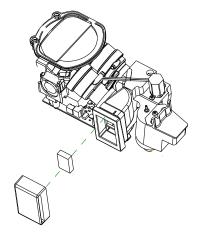


FIGURE 8-70: FILTER AND WHISPER CAP REMOVAL/INSTALLATION



8.7.2 OXYGEN BLENDER HOUSING REMOVAL

Removal

- 1. Remove the Filter/Whisper Cap. Refer to Oxygen Blender Filter/Whisper Cap Replacement Section for more details.
- 2. Remove the metal ring from the front of the Housing.
- 3. Remove the six screws securing the Manifold to the Housing.
- 4. Lift the Manifold out of the Housing.

Installation

- 1. Place the Manifold into the Housing.
- 2. Secure the Manifold to the Housing by tightening the four screws to 8 in-lbs.
- 3. Secure the two screws that connect the Mixer the to Housing by tightening the screws to 6 in-lbs.
- 4. Secure the metal ring to the assembly by tightening the screws to 6 in-lbs.
- 5. Install the Filter/Whisper Cap. Refer to Oxygen Blender Filter/Whisper Cap Replacement Section for more details.

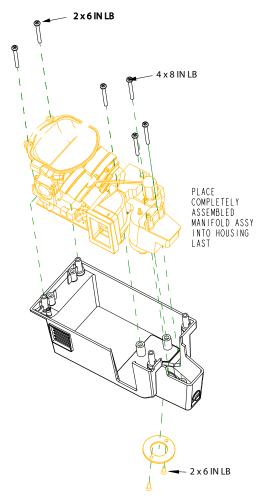


FIGURE 8-71: OXYGEN BLENDER HOUSING REMOVAL/INSTALLATION

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8.7.3 MIXING ELEMENT/FLOW ELEMENT TUBES REPLACEMENT

Removal

- Remove the Filter/Whisper Cap. Refer to Oxygen Blender Filter/Whisper Cap Replacement Section for more details.
- Remove the Oxygen Blender Housing. Refer to the Oxygen Blender Housing Removal Section for more details.
- 3. Release the Pressure Sensor Clip from the PCA.
- 4. Remove the rest of the tubing from the Flow Element Tubes.
- 5. Remove the screw securing the Mixing Element to the PCA.
- 6. Pull the Mixing element still connected to the Flow Element Tubes away from the Manifold.
- 7. Unscrew the screw securing the Flow Element Tubes to the Mixing Element.

Installation

- 1. Ensure that the two o-rings are in place at the end of the Flow Element Tubes.
- 2. Dip the end of the Flow Element Tubes in an alcohol bath and inert them into the Mixing Element.
- 3. Secure the Flow Element Tubes to the Mixing Element by securing the screw to 6 in-lbs.

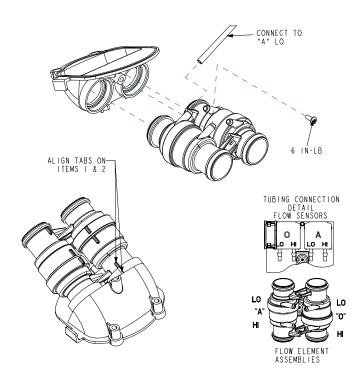


FIGURE 8-72: MIXING ELEMENT/FLOW TUBES REPLACEMENT

4. Ensure that the two o-rings are in place at the other end of the Flow Element Tubes.



5. Using a standoff, secure the Mixing Element to the PCA be securing the screw to 6 in-lbs.

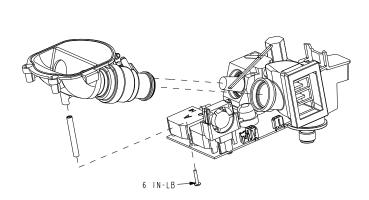


FIGURE 8-73: TUBE INSERTION AND STANDOFF PLACEMENT

- 6. Reconnect the tubing to the Flow Element Tubes and ensure it is connected properly to the PCA flow sensors.
- 7. Ensuring that the two small o-rings are in place reconnect the Pressure Sensor Clip to the PCA.
- 8. Install the Oxygen Blender Housing. Refer to Oxygen Blender Housing Removal Section for more details.
- 9. Install the Filter/Whisper Cap. Refer to Oxygen Blender Filter/Whisper Cap Replacement Section for more details.

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8.7.4 OXYGEN BLENDER PCA/PCA SPACER REPLACEMENT

Removal

- 1. Remove the Filter/Whisper Cap. Refer to Oxygen Blender Filter/Whisper Cap Replacement Section for more details.
- 2. Remove the Oxygen Blender Housing. Refer to Oxygen Blender Housing Removal Section for more details.
- 3. Remove the Mixing Element/Flow Element Tubes. Refer to Mixing Element/Flow Element Tubes Replacement Section for more details.
- 4. Disconnect the Valve connection from location P4 on the PCA.
- 5. Remove the two screws that secure the PCA to the Manifold.
- 6. Remove the screw that connect the PCA Spacer to the Manifold.

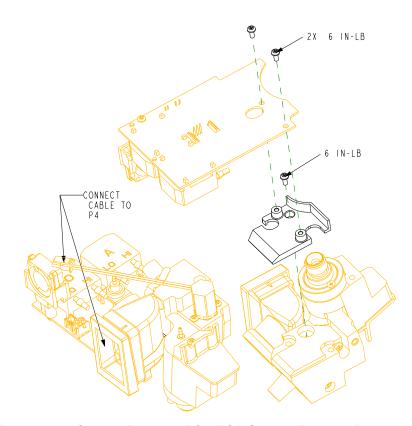


FIGURE 8-74: OXYGEN BLENDER PCA/PCA SPACER REMOVAL/INSTALLATION



Installation

- 1. Connect the PCA Spacer to the Manifold by tightening the screw to 6 in-lbs.
- 2. Connect the PCA to the Manifold be securing the two screws to 6 in-lbs.
- 3. Connect the Valve Wire to location P4 on the Main PCA.
- 4. Install the Mixing Element/Flow Element Assembly. Refer to Mixing Element/Flow Element Tubes Replacement Section for more details.
- 5. Install the Oxygen Blender Housing. Refer to Oxygen Blender Housing Removal Section for more details.
- 6. Install the Filter/Whisper Cap. Refer to Oxygen Blender Filter/Whisper Cap Replacement Section for more details.



8.7.5 OXYGEN BLENDER PRESSURE SENSOR PORT CLIP REPLACEMENT

Removal

- Remove the Filter/Whisper Cap. Refer to Oxygen Blender Filter/Whisper Cap Replacement Section for more details.
- Remove the Oxygen Blender Housing. Refer to Oxygen Blender Housing Removal Section for more details.
- 3. Remove the Mixing Element/Flow Element Tubes. Refer to Mixing Element/Flow Element Tubes Replacement Section for more details.
- Remove the Oxygen Blender PCA/PCA Spacer. Refer to Oxygen Blender PCA/PCA Spacer Replacement Section for more details.
- 5. Unlatch the Port Clip from the bottom of the Oxygen Blender PCA.
- 6. Remove the Port Clip with green tube attached and two o-rings from the Oxygen Blender PCA.

Installation

- 1. Verify the green tube is inserted flush against the mating surface on the new Port Clip.
- Insert the first new o-ring into the new Port Clip.
- 3. Insert the second new o-ring onto U24 on the Oxygen Blender PCA.
- 4. Insert the Port Clip with the first new o-ring over U24.
- 5. Latch the Port Clip onto the Oxygen Blender PCA.

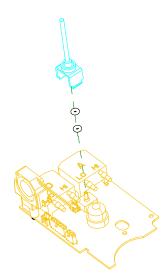


FIGURE 8-75: PORT CLIP REMOVAL/INSTALLATION

- 6. Install the Oxygen Blender PCA/PCA Spacer. Refer to Oxygen Blender PCA/PCA Spacer Replacement Section for more details.
- 7. Install the Mixing Element/Flow Element Tubes. Refer to Mixing Element/Flow Element Tubes Replacement Section for more details.
- 8. Install the Oxygen Blender Housing. Refer to Oxygen Blender Housing Removal Section for more details.
- 9. Install the Filter/Whisper Cap. Refer to Oxygen Blender Filter/Whisper Cap Replacement Section for more details.

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8.7.6 OXYGEN BLENDER FAN REPLACEMENT

Removal

- 1. Remove the Filter/Whisper Cap. Refer to Oxygen Blender Filter/Whisper Cap Replacement Section for more details.
- 2. Remove the Oxygen Blender Housing. Refer to Oxygen Blender Housing Removal Section for more details.
- 3. Disconnect the Fan wire from location P5 on the Oxygen Blender PCA.
- 4. Remove the Fan from the Fan Bracket.

Installation

1. Secure the Fan to the Fan Bracket.

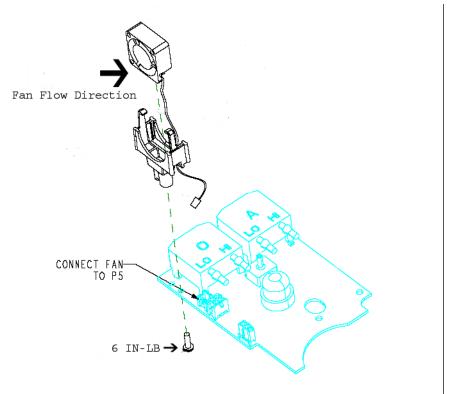


FIGURE 8-76: FAN REMOVAL/INSTALLATION

- 2. Connect the Fan Wire to location P5 on the Oxygen Blender PCA.
- 3. Install the Oxygen Blender Housing. Refer to Oxygen Blender Housing Removal Section for more details.
- 4. Install the Filter/Whisper Cap. Refer to Oxygen Blender Filter/Whisper Cap Replacement Section for more details.

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8.7.7 AIR INLET DUCT ASSEMBLY REPLACEMENT

Removal

- 1. Remove the Filter/Whisper Cap. Refer to Oxygen Blender Filter/Whisper Cap Replacement Section for more details.
- 2. Remove the Oxygen Blender Housing. Refer to Oxygen Blender Housing Removal Section for more details.
- 3. Remove the Mixing Element/Flow Element Assembly. Refer to Mixing Element/Flow Element Tubes Replacement Section for more details.
- 4. Remove the two screws that secure the Air Inlet Duct Assembly to the Manifold.

Installation

1. Connect the Air Inlet Duct Assembly to the Manifold by tightening the screws to 8 in-lbs.

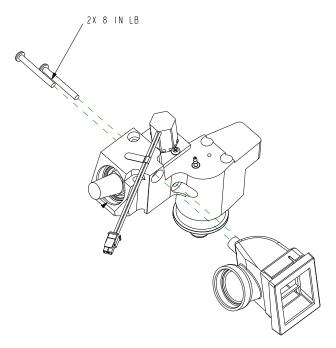


FIGURE 8-77: AIR INLET DUCT INSTALLATION

- 2. Install the Mixing Element/Flow Element Assembly. Refer to Mixing Element/Flow Element Tubes Replacement Section for more details.
- 3. Install the Oxygen Blender Housing. Refer to Oxygen Blender Housing Removal Section for more details.
- 4. Install the Filter/Whisper Cap. Refer to Oxygen Blender Filter/Whisper Cap Replacement Section for more details.



8.7.8 LOWER MANIFOLD COVER REPLACEMENT

Removal

1. Remove the screw that secures the Lower Manifold Cover to the Manifold.

Installation

1. Connect the Lower Manifold Cover to the Manifold by tightening the screw to 6 in-lbs.

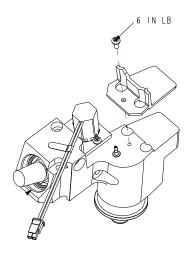


FIGURE 8-78: LOWER MANIFOLD COVER INSTALLATION

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8.7.9 Manifold Replacement

Removal

- Remove the Filter/Whisper Cap. Refer to Oxygen Blender Filter/Whisper Cap Replacement Section for more details.
- Remove the Oxygen Blender Housing. Refer to Oxygen Blender Housing Removal Section for more details.
- 3. Remove the Mixing Element/Flow Element Assembly. Refer to Mixing Element/Flow Element Tubes Replacement Section for more details.
- Remove the Oxygen Blender PCA. Refer to Oxygen Blender PCA/PCA Spacer Replacement Section for more details.
- Remove the Air Inlet Duct Assembly. Refer to the Air Inlet Duct Assembly Replacement Section for more details.
- 6. Remove the Lower Manifold Cover. Refer to the Lower Manifold Cover Replacement Section for more details.

Installation

- Install the Lower Manifold Cover. Refer to the Lower Manifold Cover Replacement Section for more details.
- Install the Air Inlet Duct Assembly. Refer to the Air Inlet Duct Assembly Replacement Section for more details.
- Install the Oxygen Blender PCA. Refer to Oxygen Blender PCA/PCA Spacer Replacement Section for more details.
- 4. Install the Mixing Element/Flow Element Assembly. Refer to Mixing Element/Flow Element Tubes Replacement Section for more details.
- 5. Install the Oxygen Blender Housing. Refer to Oxygen Blender Housing Removal Section for more details.
- 6. Install the Filter/Whisper Cap. Refer to Oxygen Blender Filter/Whisper Cap Replacement Section for more details.



8.8 PACKAGING ASSEMBLY PROCEDURE

1. Form device shipper box and seal, using 3 inch tape.



2. Form & insert the Bottom Pad and place it into shipper.



3. Place device with handle down into appropriately-sized Polybag, folding excess polybag around device.

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4. Place the bagged device into shipper.



5. Form four roll-up spacers and place them into the shipper.



6. Form the spacer pad and place it inside the shipper.





7. Form the Top pad and insert into the shipper box.



- 8. Close the shipper box and seal with 3 inch tape.
- 9. Place shipping labels on the outside of the shipper box.



CHAPTER 9: REPAIR KITS

9.0 CHAPTER OVERVIEW

This chapter illustrates the names and components for each of the repair kits for the Trilogy Ventilators. For technical assistance or replacement part ordering information, contact Respironics Product Support.

USA and Canada

Phone: 1-800-345-6443 Fax: 1-800-866-0245 Email: service@respironics.com

International

Phone: 1-724-387-4000 Fax: 1-800-387-5012

Visit Respironics Home Page on the World Wide Web at:

www.respironics.com



9.1 REPAIR KIT REFERENCE TABLE

NOTE

For kits with multiple part number listings, refer to the individual page to ensure proper ordering.

PART NUMBER(S)	REPAIR KIT NAME	PAGE IDENTIFIER
1045299	AC Power Connector Cable Kit	page 12
1045153	Active Exhalation Control Module Kit	page 24
1045295/1061268/1061269/ 1070258/1070256/1075608/ 1075638	Base Enclosure Kits	page 18
1045180	Battery Fan Kit	page 10
1045171	Battery Fan Shield Kit	page 25
1045307	Bellows Clip Kit	page 33
1045309	Blower Bellows Kit	page 6
1045302	Capacitor Kit	page 10
1045303	Capacitor/Battery Retainer Assembly Kit	page 7
1045589	DC Inlet O-ring Kit	page 45
1045300	DC Power Connector Cable Kit	page 11
1045166	Detachable Battery Connector Assembly Kit	page 22
1045167	Detachable Battery Retainer Kit	page 21
1079058	Enclosure Auxiliary Cover Kit	page 58
1045181	Enclosure Seal Kit	page 29
1045178	Exhaust Fan Assembly Kit	page 27
1067047	FAA Label Kit	page 58
1045150/1070249	Flow Sensor Assembly Kits	page 28
1045168	Flow Straightener Kit	page 27
1045200	Front Panel/Keypad LED PCA Kit	page 36
1045306	Handle Kit	page 6
1045590	Handle O-ring Kit	page 46
1045587	Inlet Air Path Assembly Kit	page 7
1045296	Interface Board Retainer Kit	page 20
1045297	Interface PCA Kit	page 34
1055806/1055957	Internal Battery Pack Kits	page 8
1045183	Keypad Kit	page 16
1045184	LCD Kit	page 39
1054951	Motor Blower Assembly Kit	page 34
1045588	O2 Inlet O-ring Kit	page 45

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PART NUMBER(S)	REPAIR KIT NAME	PAGE IDENTIFIER
1045170	Outlet Flow Path Thermistor Kit	page 26
1045298	Oxygen Connector Kit	page 11
1045163	PCA Inverter Kit	page 32
1045197	PCB Module Plate "A" Kit	page 16
1045198	PCB Module Plate "B" Kit	page 17
1045196	PCB Support for Power Management PCA Kit	page 13
1045195	PCB Support for System PCA Kit	page 12
1035443/1029330	Pollen Filter Kits	page 5
1045174	Porting Block Adaptor Kit	page 23
1045304	Power Cord Clamp Kit	page 13
1045201	Power Management PCA Kit	page 37
1045151	Power Supply PCA Kit	page 38
1058376	Preventive Maintenance Label Kit	page 47
1045149/1070251	Rear Enclosure Kits	page 31
1045403	Removable Air Path Foam Kit	page 5
1029360	Rubber Feet Kit	page 47
1051801	SD Card Kit	page 39
1045152/1070248	Sensor Board Assembly Kits	page 35
1045310/1055430	Shipping Container Kits	page 40
1045301	Speaker Hold-Down Kit	page 8
1064699	Speaker Kit	page 32
1045308	Stirring Fan Foam Kit	page 40
1045176	Stirring Fan Kit	page 23
1045177	Stirring Fan Retainer Kit	page 25
1045204	System Board to Ethernet Cable Kit	page 9
1045203	System Board to Interface PCA Cable Kit	page 9
1045164	System Board to LCD Cable Ferrite Kit	page 14
1045225	System Board to LCD Cable Kit	page 15
1045202	System Board to Sensor Board Cable Kit	page 15
1045199	System Board w/ Daughter Board Kit	page 37
1045169	Temperature Sensor O-ring Kit	page 26
1045586	Top Plate Enclosure Kit	page 44
1045154	Transition Tube Kit	page 24
1045165	Transition Tube Locator Kit	page 21
1045172	Trilogy 100 Tubing Kit	page 38
1070250	Trilogy 200, Trilogy O2, & Trilogy 202 Tubing Kit	page 57
1076003	Trilogy Base Seal Kit	page 58



PART NUMBER(S)	REPAIR KIT NAME	PAGE IDENTIFIER
1045182/1070253/1070252/ 1075607	Trilogy Front Enclosure Kits	page 30
1045175	Trilogy Hardware Kit	page 38
1070135	Trilogy O2 & Trilogy 202 Blender Plastic Threaded Cap	page 56
1054773	Trilogy O2 & Trilogy 202 Oxygen Blender Filter Duct	page 53
1054870	Trilogy O2 & Trilogy 202 Oxygen Blender Flow Element	page 48
1054872	Trilogy O2 & Trilogy 202 Oxygen Blender Housing	page 50
1054772	Trilogy O2 & Trilogy 202 Oxygen Blender Lower Manifold	page 54
1054774	Trilogy O2 & Trilogy 202 Oxygen Blender Manifold Assembly	page 56
1054871	Trilogy O2 & Trilogy 202 Oxygen Blender Mixing Element	page 49
1054867	Trilogy O2 & Trilogy 202 Oxygen Blender PCA	page 47
1054875	Trilogy O2 & Trilogy 202 Oxygen Blender PCA Spacer	page 51
1054868	Trilogy O2 & Trilogy 202 Oxygen Blender Pressure Sensor Port Clip	page 55
1054775	Trilogy O2 & Trilogy 202 Oxygen Blender Purge Fan	page 54
1054876	Trilogy O2 & Trilogy 202 Oxygen Blender Tubing Kit	page 57
1054869	Trilogy O2 & Trilogy 202 Oxygen Blender Whisper Cap	page 48
1054874	Trilogy O2 & Trilogy 202 Oxygen Blender Wire Harness	page 52
1070259	Trilogy Porting Block Adaptor Cap Kit	page 57
1045173	Tubing Elbow Kit	page 14
1045312/1045402/1058857/ 1070257/1070255/1075609/ 1075637	Warning Label Kits	page 41

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9.2 POLLEN FILTER KITS

PART NUMBER: 1035443 (1 PACK)/1029330 (2 PACK)		
Included in Kit	Tools Required	
Pollen Filter	None	



9.3 REMOVABLE AIR PATH FOAM KIT

PART NUMBER: 1045403		
Included in Kit	Tools Required	
Removable Air Path Foam	Phillips Screwdriver	





9.4 HANDLE KIT

PART NUMBER: 1045306		
Included in Kit	Tools Required	
Trilogy Handle	1/8" Hex Wrench	



9.5 BLOWER BELLOWS KIT

PART NUMBER: 1045309	
Included in Kit	Tools Required
Blower Bellow	Phillips Screwdriver



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9.6 INLET AIR PATH ASSEMBLY KIT

PART NUMBER: 1045587		
Included in Kit	Tools Required	
Inlet Air Path Assembly	Phillips Screwdriver	



9.7 CAPACITOR/BATTERY RETAINER ASSEMBLY KIT

PART NUMBER: 1045303	
Included in Kit	Tools Required
Capacitor/Battery Retainer Assembly	Phillips Screwdriver





9.8 INTERNAL BATTERY PACK KITS

PART NUMBER: 1055806		PART NUMBER: 1	1055957
Included in Kit	Tools Required	Included in Kit	Tools Required
International/Domestic Internal Battery Pack	Phillips Screwdriver	Japanese Internal Battery Pack	Phillips Screwdriver



9.9 SPEAKER HOLD-DOWN KIT

PART NUMBER: 1045301	
Included in Kit	Tools Required
Speaker Hold-Down	Phillips Screwdriver



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9.10 SYSTEM BOARD TO ETHERNET CABLE KIT

PART NUMBER: 1045204		
Included in Kit	Tools Required	
System Board to Ethernet Cable	Phillips Screwdriver	



9.11 SYSTEM BOARD TO INTERFACE PCA CABLE KIT

PART NUMBER: 1045203	
Included in Kit	Tools Required
System Board to Interface PCA Cable	Phillips Screwdriver





9.12 CAPACITOR KIT

PART NUMBER: 1045302		
Included in Kit	Tools Required	
Capacitor	Phillips Screwdriver	



9.13 BATTERY FAN KIT

PART NUMBER: 1045180		
Included in Kit	Tools Required	
Battery Fan	Phillips Screwdriver	





9.14 DC POWER CONNECTOR CABLE KIT

PART NUMBER: 1045300		
Included in Kit	Tools Required	
DC Power Connector Cable	Phillips Screwdriver 3/4" Wrench	



9.15 OXYGEN CONNECTOR KIT

PART NUMBER: 1045298	
Included in Kit	Tools Required
Oxygen Connector	Phillips Screwdriver 5/8" Wrench





9.16 AC POWER CONNECTOR CABLE KIT

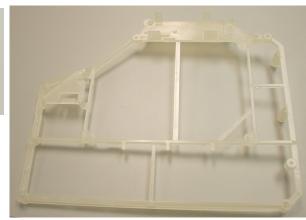
PART NUMBER: 1045299		
Included in Kit	Tools Required	
AC Power Connector Cable w/ Gasket (comes assembled)	Phillips Screwdriver 1/4" Wrench	



9.17 PCB SUPPORT FOR SYSTEM PCA KIT

PART NUMBER: 1045195		
Included in Kit	Tools Required	
PCB Support for System PCA w/ Ferrite (comes assembled)	Phillips Screwdriver 1/8" Hex Wrench	



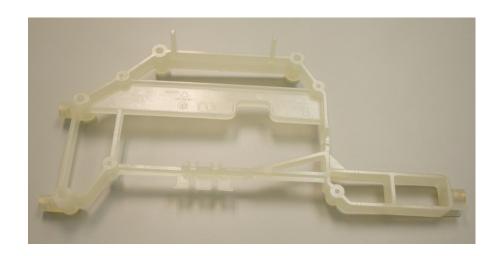


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9.18 PCB SUPPORT FOR POWER MANAGEMENT PCA KIT

PART NUMBER: 1045196	
Included in Kit	Tools Required
PCB Support for Power Management PCA	Phillips Screwdriver 1/8" Hex Wrench



9.19 POWER CORD CLAMP KIT

PART NUMBER: 1045304		
Included in Kit	Tools Required	
Power Cord Clamp	Phillips Screwdriver	





9.20 TUBING ELBOW KIT

PART NUMBER: 1045173		
Included in Kit	Tools Required	
Tubing Elbow	Phillips Screwdriver	



9.21 SYSTEM BOARD TO LCD CABLE FERRITE KIT

PART NUMBER: 1045164	
Included in Kit	Tools Required
Ferrite	Phillips Screwdriver

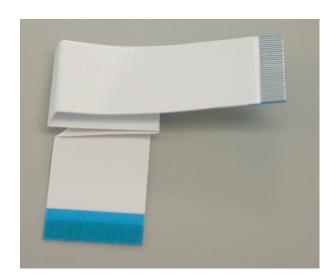


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9.22 SYSTEM BOARD TO LCD CABLE KIT

PART NUMBER: 1045225	
Included in Kit	Tools Required
System Board to LCD Cable	Phillips Screwdriver 1/8" Hex Wrench



9.23 SYSTEM BOARD TO SENSOR BOARD CABLE KIT

PART NUMBER: 1045202	
Included in Kit	Tools Required
System Board to Sensor Board Cable	Phillips Screwdriver 1/8" Hex Wrench





9.24 KEYPAD KIT

PART NUMBER: 1045183	
Included in Kit	Tools Required
Keypad	Phillips Screwdriver 1/8" Hex Wrench



9.25 PCB MODULE PLATE "A" KIT

PART NUMBER: 1045197		
Included in Kit	Tools Required	
PCB Module Plate "A"	Phillips Screwdriver 1/8" Hex Wrench	

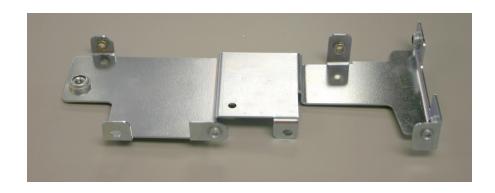


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9.26 PCB MODULE PLATE "B" KIT

PART NUMBER: 1045198	
Included in Kit	Tools Required
PCB Module Plate "B"	Phillips Screwdriver 1/8" Hex Wrench





9.27 BASE ENCLOSURE KITS

PART NUMBER: 1045295	
Included in Kit	Tools Required
Base Enclosure Assembly Clear Overlay (x2) Trilogy 100/Trilogy 200 Warning Label - International	Phillips Screwdriver

PART NUMBER: 1061269	
Included in Kit	Tools Required
Base Enclosure Assembly Clear Overlay (x2) Trilogy 100/Trilogy 200 Warning Label - Domestic	Phillips Screwdriver

PART NUMBER: 1061268	
Included in Kit	Tools Required
Base Enclosure Assembly Clear Overlay (x2) Trilogy 100/Trilogy 200 Warning Label - Japanese	Phillips Screwdriver

PART NUMBER: 1070258	
Included in Kit	Tools Required
Base Enclosure Assembly Clear Overlay (x2) Trilogy O ₂ Warning Label - Japanese	Phillips Screwdriver

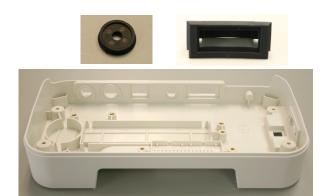
PART NUMBER: 1070256	
Included in Kit	Tools Required
Base Enclosure Assembly Clear Overlay (x2) Trilogy 100/Trilogy 200 Warning Label - Latin America	Phillips Screwdriver

PART NUMBER: 1075638	
Included in Kit	Tools Required
Base Enclosure Assembly Clear Overlay (x2) Trilogy 202 Warning Label - International	Phillips Screwdriver

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PART NUMBER: 1075608	
Included in Kit	Tools Required
Base Enclosure Assembly Clear Overlay (x2) Trilogy 202 Warning Label - Domestic	Phillips Screwdriver





9.28 INTERFACE BOARD RETAINER KIT

PART NUMBER: 1045296	
Included in Kit	Tools Required
Interface Board Retainer	Phillips Screwdriver



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9.29 TRANSITION TUBE LOCATOR KIT

PART NUMBER: 1045165	
Included in Kit	Tools Required
Transition Tube Locator	Phillips Screwdriver 1/8" Hex Wrench



9.30 DETACHABLE BATTERY RETAINER KIT

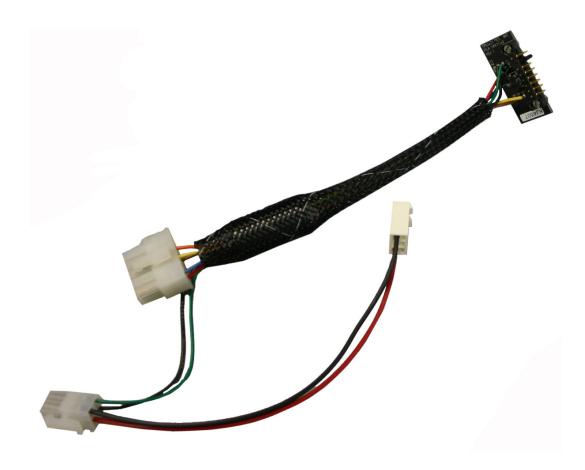
PART NUMBER: 1045167	
Included in Kit	Tools Required
Detachable Battery Retainer	Phillips Screwdriver 1/8" Hex Wrench





9.31 DETACHABLE BATTERY CONNECTOR ASSEMBLY KIT

PART NUMBER: 1045166	
Included in Kit	Tools Required
Detachable Battery Connector Assembly	Phillips Screwdriver 1/8" Hex Wrench



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9.32 STIRRING FAN KIT

PART NUMBER: 1045176	
Included in Kit	Tools Required
Stirring Fan	Phillips Screwdriver 1/8" Hex Wrench



9.33 PORTING BLOCK ADAPTOR KIT

PART NUMBER: 1045174	
Included in Kit	Tools Required
Porting Block Adaptor	Phillips Screwdriver 1/8" Hex Wrench





9.34 TRANSITION TUBE KIT

PART NUMBER: 1045154	
Included in Kit	Tools Required
Transition Tube	Phillips Screwdriver 1/8" Hex Wrench



9.35 ACTIVE EXHALATION CONTROL MODULE KIT

PART NUMBER: 1045153	
Included in Kit	Tools Required
Active Exhalation Control Module O-rings (x2)	Phillips Screwdriver 1/8" Hex Wrench



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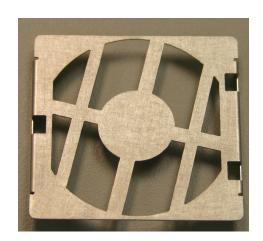
9.36 STIRRING FAN RETAINER KIT

PART NUMBER: 1045177	
Included in Kit	Tools Required
Stirring Fan Retainer Stirring Fan Foam	Phillips Screwdriver 1/8" Hex Wrench



9.37 BATTERY FAN SHIELD KIT

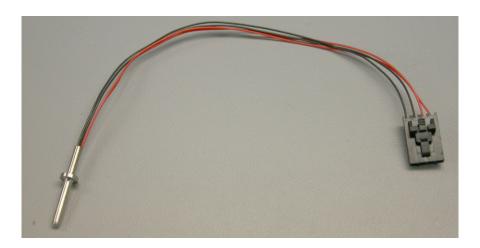
PART NUMBER: 1045171	
Included in Kit	Tools Required
Battery Fan Shield	Phillips Screwdriver





9.38 OUTLET FLOW PATH THERMISTOR KIT

PART NUMBER: 1045170	
Included in Kit	Tools Required
Outlet Flow Path Thermistor	Phillips Screwdriver 1/8" Hex Wrench



9.39 TEMPERATURE SENSOR O-RING KIT

PART NUMBER: 1045169	
Included in Kit	Tools Required
Temperature Sensor O-ring	Phillips Screwdriver 1/8" Hex Wrench



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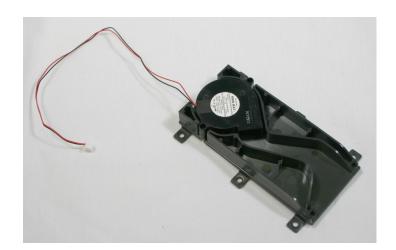
9.40 FLOW STRAIGHTENER KIT

PART NUMBER: 1045168	
Included in Kit	Tools Required
Flow Straightener	Phillips Screwdriver 1/8" Hex Wrench Chamfer tool M100311



9.41 EXHAUST FAN ASSEMBLY KIT

PART NUMBER: 1045178	
Included in Kit	Tools Required
Exhaust Fan Assembly	Phillips Screwdriver 1/8" Hex Wrench





9.42 FLOW SENSOR ASSEMBLY KITS

PART NUMBER: 1045150	
Included in Kit	Tools Required
Trilogy 100 Flow Sensor Assembly	Phillips Screwdriver 1/8" Hex Wrench



PART NUMBER: 1070249	
Included in Kit	Tools Required
Trilogy 200/Trilogy 202/Trilogy O ₂ Flow Sensor Assembly	Phillips Screwdriver 1/8" Hex Wrench



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9.43 ENCLOSURE SEAL KIT

PART NUMBER: 1045181	
Included in Kit	Tools Required
Enclosure Seal	Phillips Screwdriver 1/8" Hex Wrench





9.44 TRILOGY FRONT ENCLOSURE KITS

PART NUMBER: 1045182	
Included in Kit	Tools Required
Trilogy 100 Front Enclosure	Phillips Screwdriver 1/8" Hex Wrench

PART NUMBER: 1070253	
Included in Kit	Tools Required
Trilogy O ₂ Front Enclosure	Phillips Screwdriver 1/8" Hex Wrench

PART NUMBER: 1070252	
Included in Kit	Tools Required
Trilogy 200 Front Enclosure	Phillips Screwdriver 1/8" Hex Wrench

PART NUMBER: 1075607	
Included in Kit	Tools Required
Trilogy 202 Front Enclosure	Phillips Screwdriver 1/8" Hex Wrench



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9.45 REAR ENCLOSURE KITS

PART NUMBER: 1045149	
Included in Kit	Tools Required
Trilogy 100/Trilogy 200 Rear Enclosure	Phillips Screwdriver 1/8" Hex Wrench

PART NUMBER: 1070251	
Included in Kit	Tools Required
Trilogy 202/Trilogy O ₂ Rear Enclosure (contains hole for Oxygen Blending Module connection)	Phillips Screwdriver 1/8" Hex Wrench





9.46 PCA INVERTER KIT

PART NUMBER: 1045163	
Included in Kit	Tools Required
PCA Inverter	Phillips Screwdriver 1/8" Hex Wrench



9.47 SPEAKER KIT

PART NUMBER: 1064699	
Included in Kit	Tools Required
Speaker	Phillips Screwdriver



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9.48 BELLOWS CLIP KIT

PART NUMBER: 1045307	
Included in Kit	Tools Required
Bellows Clip	Phillips Screwdriver





9.49 MOTOR BLOWER ASSEMBLY KIT

PART NUMBER: 1054951	
Included in Kit	Tools Required
Motor Blower Assembly w/ Isolation Assembly	Phillips Screwdriver 1/8" Hex Wrench



9.50 INTERFACE PCA KIT

PART NUMBER: 1045297	
Included in Kit	Tools Required
Interface PCA	Phillips Screwdriver



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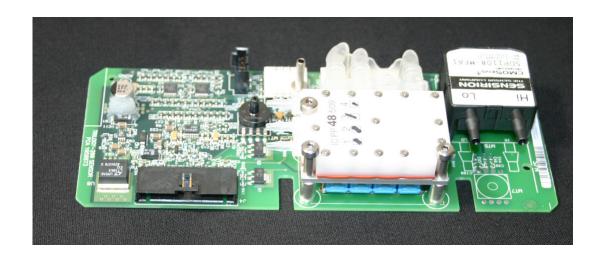


9.51 SENSOR BOARD ASSEMBLY KITS

PART NUMBER: 1045152	
Included in Kit	Tools Required
Trilogy 100 Sensor Board Assembly	Phillips Screwdriver 1/8" Hex Wrench



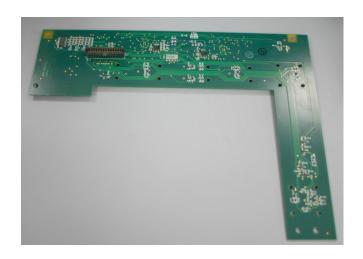
PART NUMBER: 1070248	
Included in Kit	Tools Required
Trilogy 200/Trilogy 202/Trilogy O ₂ Sensor Board Assembly	Phillips Screwdriver 1/8" Hex Wrench





9.52 FRONT PANEL/KEYPAD LED PCA KIT

PART NUMBER: 1045200	
Included in Kit	Tools Required
Front Panel/Keypad LED PCA	Phillips Screwdriver 1/8" Hex Wrench



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9.53 SYSTEM BOARD W/ DAUGHTER BOARD KIT

PART NUMBER: 1045199	
Included in Kit	Tools Required
System Board w/ Daughter Board	Phillips Screwdriver 1/8" Hex Wrench



9.54 POWER MANAGEMENT PCA KIT

PART NUMBER: 1045201	
Included in Kit	Tools Required
Power Management PCA	Phillips Screwdriver 1/8" Hex Wrench





9.55 POWER SUPPLY PCA KIT

PART NUMBER: 1045151	
Included in Kit	Tools Required
Power Supply PCA	Phillips Screwdriver 1/8" Hex Wrench



9.56 TRILOGY 100 TUBING KIT

PART NUMBER: 1045172	
Included in Kit	Tools Required
Trilogy 100 Tubing	Phillips Screwdriver 1/8" Hex Wrench

9.57 TRILOGY HARDWARE KIT

PART NUMBER: 1045175	
Included in Kit	Tools Required
Trilogy Hardware Kit	

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9.58 LCD KIT

PART NUMBER: 1045184	
Included in Kit	Tools Required
Trilogy Color 5.7 Inch LCD	Phillips Screwdriver 1/8" Hex Wrench



9.59 SD CARD KIT

PART NUMBER: 1051801	
Included in Kit	Tools Required
1 GB SD Card	None





9.60 SHIPPING CONTAINER KITS

PART NUMBER: 1045310 (1 PACK) / 1055430 (10 PACK)	
Included in Kit	Tools Required
Trilogy Shipping Container	

9.61 STIRRING FAN FOAM KIT

PART NUMBER: 1045308	
Included in Kit	Tools Required
Stirring Fan Foam	

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9.62 WARNING LABEL KITS

PART NUMBER: 1045312	
Included in Kit	Tools Required
Trilogy 100/Trilogy 200 Warning Label (International)	N/A



PART NUMBER: 1045402	
Included in Kit	Tools Required
Trilogy 100/Trilogy 200 Warning Label (Japanese)	N/A





PART NUMBER: 1058857	
Included in Kit	Tools Required
Trilogy 100/Trilogy 200 Warning Label (Domestic)	N/A



PART NUMBER: 1070257	
Included in Kit	Tools Required
Trilogy O ₂ Warning Label	N/A

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PART NUMBER: 1070255	
Included in Kit	Tools Required
Trilogy 100/Trilogy 200 Warning Label (Latin America)	N/A





PART NUMBER: 1075609	
Included in Kit	Tools Required
Trilogy 202 Warning Label (Domestic)	N/A

PART NUMBER: 1075637	
Included in Kit	Tools Required
Trilogy 202 Warning Label (International)	N/A

9.63 TOP PLATE ENCLOSURE KIT

PART NUMBER: 1045586	
Included in Kit	Tools Required
Top Plate	Phillips Screwdriver 1/8" Hex Wrench



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9.64 O₂ Inlet O-RING KIT

PART NUMBER: 1045588	
Included in Kit	Tools Required
O ₂ Inlet O-ring	Phillips Screwdriver 1/8" Hex Wrench 5/8" Wrench



9.65 DC INLET O-RING KIT

PART NUMBER: 1045589	
Included in Kit	Tools Required
DC Inlet O-ring	Phillips Screwdriver 1/8" Hex Wrench 3/4" Wrench

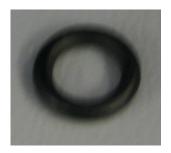


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9.66 HANDLE O-RING KIT

PART NUMBER: 1045590	
Included in Kit	Tools Required
Handle O-ring	1/8" Hex Wrench



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9.67 RUBBER FEET KIT

PART NUMBER: 1029360	
Included in Kit	Tools Required
Rubber Feet	None

9.68 PREVENTIVE MAINTENANCE LABEL KIT

PART NUMBER: 1058376	
Included in Kit	Tools Required
Preventive Maintenance Label (10 pack)	None

9.69 TRILOGY O_2 & TRILOGY 202 OXYGEN BLENDER PCA

PART NUMBER: 1054867	
Included in Kit	Tools Required
Oxygen Blender PCA	None

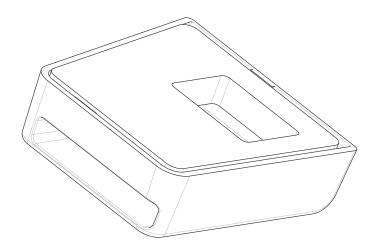


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9.70 TRILOGY O_2 & TRILOGY 202 OXYGEN BLENDER WHISPER CAP

PART NUMBER: 1054869	
Included in Kit	Tools Required
Oxygen Blender Whisper Cap	None



9.71 TRILOGY O_2 & TRILOGY 202 OXYGEN BLENDER FLOW ELEMENT

PART NUMBER: 1054870	
Included in Kit	Tools Required
Oxygen Blender Flow Element	None



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9.72 TRILOGY O_2 & TRILOGY 202 OXYGEN BLENDER MIXING ELEMENT

PART NUMBER: 1054871	
Included in Kit	Tools Required
Oxygen Blender Mixing Element	None



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9.73 TRILOGY O_2 & TRILOGY 202 OXYGEN BLENDER HOUSING

PART NUMBER: 1054872	
Included in Kit	Tools Required
Oxygen Blender Housing	None



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9.74 TRILOGY O_2 & TRILOGY 202 OXYGEN BLENDER PCA SPACER

PART NUMBER: 1054875	
Included in Kit	Tools Required
Oxygen Blender PCA Spacer	None



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9.75 TRILOGY O₂ & TRILOGY 202 OXYGEN BLENDER WIRE HARNESS

PART NUMBER: 1054874	
Included in Kit	Tools Required
Oxygen Blender Wire Harness	None



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9.76 TRILOGY O_2 & TRILOGY 202 OXYGEN BLENDER FILTER DUCT

PART NUMBER: 1054773	
Included in Kit	Tools Required
Oxygen Blender Filter Duct	None



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9.77 TRILOGY O_2 & TRILOGY 202 OXYGEN BLENDER PURGE FAN

PART NUMBER: 1054775	
Included in Kit	Tools Required
Oxygen Blender Purge Fan	None



9.78 TRILOGY O_2 & TRILOGY 202 OXYGEN BLENDER LOWER MANIFOLD

PART NUMBER: 1054772	
Included in Kit	Tools Required
Oxygen Blender Lower Manifold	None

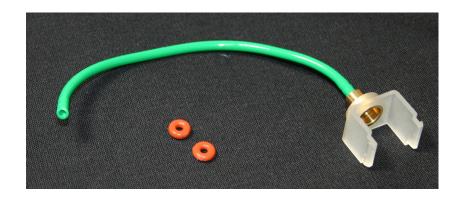


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9.79 TRILOGY O_2 & TRILOGY 202 OXYGEN BLENDER PRESSURE SENSOR PORT CLIP

PART NUMBER: 1054868	
Included in Kit	Tools Required
Oxygen Blender Pressure Sensor Port Clip	None

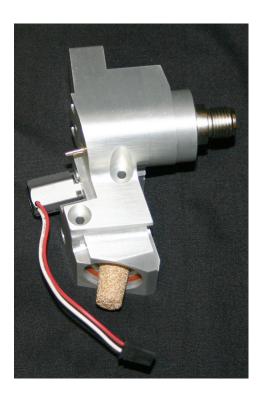


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9.80 TRILOGY O_2 & TRILOGY 202 OXYGEN BLENDER MANIFOLD ASSEMBLY

PART NUMBER: 1054774	
Included in Kit	Tools Required
Oxygen Blender Manifold Assembly	None



9.81 TRILOGY O_2 & TRILOGY 202 BLENDER PLASTIC THREADED CAP

PART NUMBER: 1070135	
Included in Kit	Tools Required
Oxygen Blender Plastic Threaded Cap	None

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9.82 Trilogy 200, Trilogy O_2 , & Trilogy 202 Tubing Kit

PART NUMBER: 1070250	
Included in Kit	Tools Required
Trilogy 200 / Trilogy O ₂ / Trilogy 202 Tubing	None

9.83 TRILOGY O_2 & TRILOGY 202 OXYGEN BLENDER TUBING KIT

PART NUMBER: 1054876	
Included in Kit	Tools Required
Oxygen Blender Tubing	None

9.84 TRILOGY PORTING BLOCK ADAPTOR CAP KIT

PART NUMBER: 1070259	
Included in Kit	Tools Required
Porting Block Adaptor Cap	None



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9.85 FAA LABEL KIT

PART NUMBER: 1067047	
Included in Kit	Tools Required
FAA Label (Qty: 10)	None



9.86 TRILOGY BASE SEAL KIT

PART NUMBER: 1076003	
Included in Kit	Tools Required
Base Enclosure Seals	None

9.87 ENCLOSURE AUXILIARY COVER KIT

PART NUMBER: 1079058	
Included in Kit	Tools Required
Enclosure Auxiliary Cover	None

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CHAPTER 10: TRILOGY TESTING & CALIBRATION

10.0 CHAPTER OVERVIEW

This chapter provides the necessary performance, service, and safety (optional) testing procedures. The intervals for the specific tests are listed in the testing procedure sections below.

10.1 TRILOGY 100 CHECKOUT PROCEDURE

This test procedure should be performed prior to connecting the device to a patient or in between patient usage. Test both the active and passive circuits if you want to do a complete checkout on the device. The tests should be performed as described in order to verify proper operation of the device.

NOTE

The actual circuit configuration to be used on the patient should be used to perform the system checkout.

10.1.1 Tools Required

- Active Exhalation Porting Block with PAP (RI p/n 1054670)
- Passive Exhalation Porting Block (RI p/n 1040372)
- Active Exhalation Device with PAP (RI p/n 1053716)
- Whisper Swivel II (RI p/n 332113)
- Test Lung (RI p/n 1021671)
- Small Flat Head Screwdriver

WARNING

If you notice any unexplained changes in the performance of the device, if it is making unusual sounds, if the device or detachable battery are dropped, if water is spilled into the enclosure or if the enclosure is cracked or broken, discontinue use and contact Respironics or an authorized service center for service.

10.1.2 VISUAL INSPECTION

- 1. Verify that the enclosure is not broken and that all applicable screws are in place.
- Verify that the device handle, SD Card door, and Detachable Battery are secure and in good working order.
- 3. Verify that the rubber feet are on the bottom of the device.

10.1.3 INITIAL SETUP

- 1. Connect the power cord to the device and then to an AC outlet.
- 2. Attach the test lung to the patient connection end of the desired circuit (Active PAP or Passive).
- 3. Access the Setup Screen. Reference the System Setup section for more information.

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10.1.4 SETTING AND ALARMS TESTS

Complete the following steps to set up the settings and alarms tests.

SETUP

Settings and Alarms Menu

Modify the settings in the Setting and Alarms menu to match those shown below in Table 10-1. If necessary refer to the System Setup Section for instructions on modifying ventilator settings.

TABLE 10-1: VENTILATOR SETTINGS IN THE SETTING AND ALARMS MENU

SETTING	VALUE
Dual Prescription	Off
Circuit Type	Active PAP or Passive
Therapy Mode	S/T
AVAPS (passive circuit only)	Off
IPAP	20 pressure units
EPAP	4 pressure units
Breath Rate	12 BPM
Inspiratory Time	1.6 seconds
Trigger Type (passive circuit)	Auto-Trak
Flow Trigger Sensitivity (active PAP circuit)	6.0 l/min
Flow Cycle Sensitivity (active PAP circuit)	20%
Rise Time	1
Ramp Length	Off
All other alarms	Off

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Options Menu

Modify the settings in the Options menu to match those shown below in Table 10-2.

TABLE 10-2: VENTILATOR SETTING IN THE OPTIONS MENU

SETTING	VALUE
Menu Access	Full
Detailed View	On
All other settings	Discretionary

Turn Device Power On

Press the Start/Stop button on the front of the ventilator. The system will begin operating using the defined ventilation settings.

VERIFY THE HIGH TIDAL VOLUME ALARM

This procedure verifies that the High Tidal Volume alarm is working properly. For passive circuits, this will verify the High Vte alarm. For active with PAP circuits, this will verify the High Vti alarm. It assumes that you have attached the test lung, verified the ventilator settings, and turned on ventilator power.

Change Alarm Ventilator Setting

Modify the High Tidal Volume alarm setting to match the one shown below in Table 10-3.

TABLE 10-3: HIGH TIDAL VOLUME ALARM SETTING

SETTING	VALUE
High Vte/High Vti	50 ml

Verify the Alarm

NOTE

Do not use the "Reset" button to manually reset the alarm. Instead, use the "Modify" button to change ventilator settings. This applies to all tests.

Wait up to 40 seconds and verify the following alarm signals:

- The High Priority audible indicator sounds
- A red light flashes on the Alarm Indicator/Audio Pause button
- The High Tidal Volume alarm condition appears on the screen, highlighted in red

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Modify Ventilator Alarm Setting

Modify the High Tidal Volume alarm setting to match the one shown below in Table 10-4.

TABLE 10-4: RESET HIGH TIDAL VOLUME ALARM SETTING

SETTING	VALUE
High Vte/Vti	500 ml

Verify Reset

Wait 40 seconds and verify the following auto-reset conditions:

- The High Priority audible indicator has stopped sounding
- The red light on the Alarm Indicator/Audio Pause button has stopped flashing

Restore Ventilator Settings

Modify the ventilator setting and change the following value shown in Table 10-5.

TABLE 10-5: RESTORE VENTILATOR SETTINGS

SETTING	VALUE
High Vte/Vti	Off

VERIFY THE LOW TIDAL VOLUME ALARM

This procedure verifies that the Low Tidal Volume alarm is working properly. For passive circuits, this will verify the Low Vte alarm. For active with PAP circuits, this will verify the Low Vti alarm. It assumes that you have attached the test lung, verified the ventilator settings, and turned on ventilator power.

Change Alarm Ventilator Setting

Modify the Low Tidal Volume alarm setting to match the one shown below in Table 10-6.

TABLE 10-6: LOW TIDAL VOLUME ALARM SETTING

SETTING	VALUE
Low Vte/Vti	500 ml

Verify the Alarm

Wait up to 40 seconds and verify the following alarm signals:

- The High Priority audible indicator sounds
- A red light flashes on the Alarm Indicator/Audio Pause button
- The Low Tidal Volume alarm condition appears on the screen, highlighted in red

Modify Ventilator Alarm Settings



Modify the Low Tidal Volume alarm setting to match the one shown below in Table 10-7.

TABLE 10-7: RESET LOW TIDAL VOLUME ALARM SETTING

SETTING	VALUE
Low Vte/Vti	50 ml

Verify Reset

Wait 40 seconds and verify the following auto-reset conditions:

- · The High Priority audible indicator has stopped sounding
- The red light on the Alarm Indicator/Audio Pause button has stopped flashing

Restore Ventilator Settings

Modify the ventilator setting and change the following value shown in Table 10-8.

TABLE 10-8: RESTORE VENTILATOR SETTINGS

SETTING	VALUE
Low Vte/Vti	Off

VERIFY THE CIRCUIT DISCONNECT ALARM

This procedure verifies that the Circuit Disconnect alarm is working properly. It assumes that you have attached the test lung, verified ventilator settings, and turned on ventilator power.

Change Circuit Disconnect Ventilator Setting

Modify the Circuit Disconnect ventilator settings to match the one shown below in Table 10-9.

TABLE 10-9: VENTILATOR SETTINGS

SETTING	VALUE
Circuit Disconnect	10 Seconds

Disconnect Test Lung

Disconnect the test lung from the circuit.

NOTE
The Low Inspiratory Pressure Alarm may also be detected.

Verify the Alarm

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Wait approximately 10 seconds and verify the following alarm signals:

- The High Priority audible indicator sounds
- A red light flashes on the Alarm Indicator/Audio Pause button
- The Circuit Disconnect alarm condition appears on the screen, highlighted in red

Reconnect Test Lung

Reconnect the test lung to the circuit.

Verify Reset

Wait at least 40 seconds and verify the following auto-reset conditions:

- The High Priority audible indicator has stopped sounding
- The red light on the Alarm Indicator/Audio Pause button has stopped flashing

Restore Ventilator Settings

Modify the ventilator setting and change the following value shown in Table 10-10.

TABLE 10-10: RESTORE VENTILATOR SETTINGS

SETTING	VALUE
Circuit Disconnect	Off

VERIFY THE HIGH INSPIRATORY PRESSURE ALARM

This procedure verifies that the High Inspiratory Pressure alarm is working properly. It assumes that you have attached the test lung, verified ventilator settings, and turned on ventilator power.

Change Ventilator Settings

Modify the ventilator settings and change the following values shown below in Table 10-11.

TABLE 10-11: VENTILATOR SETTINGS

SETTING	VALUE
Mode	CV
Tidal Volume	500 ml
Breath Rate	12 BPM
Inspiratory Time	1.0 seconds
Flow Pattern	Ramp
PEEP	4 pressure units
Sigh	Off
Circuit Disconnect	Off

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TABLE 10-11: VENTILATOR SETTINGS

SETTING	VALUE
Low Inspiratory Pressure	6 pressure units
High Inspiratory Pressure	10 pressure unit
Apnea	Off
All other alarms	Off

Verify the Alarm

Wait up to 40 seconds and verify the following alarm signals:

- The Medium Priority audible indicator sounds
- A yellow light flashes on the Alarm Indicator/Audio Pause button
- The High Inspiratory Pressure alarm condition appears on the screen, highlighted in yellow

NOTE

If this alarm is not reset within 3 occurrences, the alarm is elevated to High Priority, and the High Priority Indicators occur.

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Modify Ventilator Alarm Settings

Modify the High Inspiratory Pressure setting to match the one shown in Table 10-12.

TABLE 10-12: RESET HIGH INSPIRATORY PRESSURE ALARM

SETTING	VALUE
High Inspiratory Pressure	60 pressure units

Verify Reset

Wait 40 seconds and verify the following auto-reset conditions:

- The Medium Priority audible indicator has stopped sounding
- The yellow light on the Alarm Indicator/Audio Pause button has stopped flashing

VERIFY THE LOW INSPIRATORY PRESSURE ALARM

This procedure verifies that the Low Inspiratory Pressure alarm is working properly. It assumes that you have attached the test lung, verified ventilator settings, and turned on ventilator power.

Change Ventilator Settings

Modify the ventilator settings and change the following values shown below in Table 10-13.

TABLE 10-13: VENTILATOR SETTINGS

SETTING	VALUE
Mode	CV
Tidal Volume	500 ml
Breath Rate	12 BPM
Inspiratory Time	1.0 seconds
Flow Pattern	Ramp
PEEP	4 pressure units
Sigh	Off
Circuit Disconnect	Off
Low Inspiratory Pressure	40 pressure units
High Inspiratory Pressure	60 pressure units
Apnea	Off
All other alarms	Off

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Verify the Alarm

Wait up to 40 seconds and verify the following alarm signals:

- The High Priority audible indicator sounds
- A red light flashes on the Alarm Indicator /Audible Pause button
- The Low Inspiratory Pressure alarm condition appears on the screen, highlighted in red

Modify Ventilator Alarm Settings

Modify the Low Inspiratory Pressure setting to match the one shown below in Table 10-14.

TABLE 10-14: RESET LOW INSPIRATORY PRESSURE ALARM SETTING

SETTING	VALUE
Low Inspiratory Pressure	6 pressure units

Verify Reset

Wait 40 seconds and verify the following auto-reset conditions:

- The High Priority Indicator has stopped sounding
- The red light on the Audio Pause/Alarm Indicator button has stopped flashing

10.1.5 BATTERY FUNCTION VERIFICATION

Make sure the batteries are functioning properly and fully charged before usage.

VERIFY THE DETACHABLE AND INTERNAL (LITHIUM-ION) BATTERIES FUNCTION

- 1. Connect AC Power to the device and verify that the green AC LED on the front panel is lit.
- 2. Verify that the detachable battery is properly installed.
- 3. Turn the device on and verify that both the detachable and internal battery symbols appear on the display. Verify if either battery is less than fully charged (the charge symbol will display on the respective battery.)
- 4. Disconnect the AC Power source from the device.
 - Verify that the AC Power Disconnected alarm message appears on the display and the green AC LED is not lit. Press Reset.
 - Verify that the detachable battery symbol shows the level of charge noted in the previous step and that the device continues to operate.
 - Verify that the detachable battery symbol has a black box around it to indicate that it is in use.
- 5. Disconnect the detachable battery pack from the device.
 - Verify that the Detach Batt Disconnected alarm message appears on the display. Press Reset.
 - Verify that the internal battery symbol shows the same level of charge as noted above and the device continues to operate.
 - Verify that the internal battery symbol has a black box around it to indicate that it is in use.
- 6. Reconnect the Detachable Battery and AC Power source.



VERIFY THE EXTERNAL BATTERY FUNCTION (OPTIONAL)

- 1. Connect AC Power to the device and verify that the green AC LED is lit.
- 2. Connect the external battery cable to the external battery and to the ventilator.
- 3. Verify that the external battery symbol is shown on the display and some level of charge is present.
- 4. Disconnect the AC Power source from the device.
 - Verify that the AC Power Disconnected alarm message appears on the display and the green AC LED is not lit. Press Reset.
 - Verify that the external battery symbol shows the level of charge as noted in the previous step and the device continues to operate.
 - Verify that the external battery symbol has a black box around it to indicate that it is in use.
- 5. Reconnect the AC Power source.

ALARM AND EVENT LOG CLEAN-UP

- 1. In the Setup Menu, select Alarm Log.
- 2. Press Clear to clear the log file.
- 3. Press Yes to confirm.
- 4. Press Finish to complete.
- 5. In the Setup Menu, select **Event Log**.
- 6. Press Clear to clear the log file.
- 7. Press Yes to confirm.
- 8. Press Finish to complete.

RESULTS

All portions of this checkout procedure should be completed prior to connection to the patient. If any of the tests fail to complete as indicated, if possible, correct the error, clear the alarm and resume testing.

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10.2 TRILOGY 100 CHECKOUT PROCEDURE DATA SHEET

10.2.1 VISUAL INSPECTION

Damaged Parts?	YES	NO

10.2.2 SETTING & ALARM TESTS

High Tidal Volume Alarm Setting?	PASS	FAIL
Low Tidal Volume	PASS	FAIL
Alarm Setting?		
		•
Circuit Disconnect	PASS	FAIL
Alarm Setting?		
	1	
High Inspiratory	PASS	FAIL
Pressure Alarm Setting?		
Low Inspiratory	PASS	FAIL
Pressure Alarm Setting?		
	•	
Battery Function	PASS	FAIL
Verification?		

Signature:		 	
Date:	· · · · · · · · · · · · · · · · · · ·	 	
Serial Number:			



10.3 TRILOGY 200 CHECKOUT PROCEDURE

This chapter details the test procedures that should be performed by the clinician prior to connecting the device to the patient. Test the Active PAP, Active Flow, and Passive circuit types if you want to do a complete checkout on the device. The tests should be performed as described in order to verify proper operation of the device. Some of the procedures in this chapter require you to change settings on the device.

NOTE

The actual circuit configuration to be used on the patient should be used to perform the system checkout.

10.3.1 Tools Required

- Universal Porting Block
- Active PAP Exhalation Device
- Active Flow Exhalation Device with Flow Sensor
- Trilogy Universal Active PAP Tube Adapter
- Whisper Swivel II
- Test Lung
- Small Flat Head Screwdriver

10.3.2 VISUAL INSPECTION

- 1. Verify that the enclosure is not broken and that all applicable screws are in place.
- 2. Verify that the device handle, SD Card door, and detachable battery are secure and in good working order.
- Verify that the rubber feet are on the bottom of the device.

10.3.3 INITIAL SETUP

- 1. Connect the power cord to the device and then to an AC outlet.
- 2. Attach the test lung to the patient connection end of the desired circuit (Active PAP, Active Flow, or Passive).
- 3. Follow the instructions in System Setup Section to access the Setup Screen.

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10.3.4 SETTINGS AND ALARM TESTS

Complete the following steps to set up the settings and alarms tests.

SETUP

Settings And Alarms Menu

Modify the settings in the Settings and Alarms menu to match those shown below.

SETTING	VALUE
Dual Prescription	Off
Circuit Type	Active PAP, Active Flow or Passive
Therapy Mode	S/T
AVAPS (passive circuit only)	Off
IPAP	20 pressure units
EPAP	4 pressure units
Breath Rate	12 BPM
Inspiratory Time	1.6 seconds
Trigger Type (passive circuit)	Auto-Trak
Flow Trigger Sensitivity (active PAP circuit)	6.0 l/min
Leak Compensation (Active Flow circuit type	On
Flow Cycle Sensitivity (Active PAP or Active Flow circuit type)	20%
Rise Time	1
Ramp Length	Off
All other alarms	Off

Options Menu

Modify the settings in the Options menu to match those shown below.

SETTING	VALUE
Menu Access	Full
Detailed View	On
All other settings	Discretionary



Turn Device Power On

Press the Start/Stop button on the front of the ventilator. The system will begin operating using the defined ventilation settings.

VERIFY THE HIGH TIDAL VOLUME ALARM

NOTE

Do not use the "reset" button to manually reset the alarm. Instead, use the "Modify" button to change ventilator settings. This note applies to all tests.

This procedure verifies that the High Tidal Volume alarm is working properly. For Passive and Active Flow circuits, this will verify the High Vte alarm. For Active PAP circuits, this will verify the High Vti alarm. It assumes that you have attached the test lung, verified the ventilator settings, and turned on ventilator power as described in the Initial Setup section.

Change Alarm Ventilator Setting

Modify the High Tidal Volume alarm setting to match the one shown below.

SETTING	VALUE
High Vte/High Vti	50 ml

Verify the Alarm

Wait up to 40 seconds and verify the following alarm signals:

- The High Priority audible indicator sounds
- A red light flashes on the Alarm Indicator/Audio Pause button
- The High Tidal Volume alarm condition appears on the screen, highlighted in red

Modify Ventilator Alarm Settings

Modify the High Tidal Volume alarm setting to match the one shown below.

SETTING	VALUE
High Vte/High Vti	500 ml

Verify Reset

Wait 40 seconds and verify the following auto-reset conditions:

- The High Priority audible indicator has stopped sounding
- The red light on the Alarm Indicator/Audio Pause button has stopped flashing



Restore Ventilator Settings

Modify the ventilator settings and change the following value shown below.

SETTING	VALUE
High Vte/High Vti	Off

VERIFY THE LOW TIDAL VOLUME ALARM

This procedure verifies that the Low Tidal Volume alarm is working properly. For Passive and Active Flow circuits, this will verify the Low Vte alarm. For Active PAP circuits, this will verify the Low Vti alarm. It assumes that you have attached the test lung, verified the ventilator settings, and turned on ventilator power as described in the Initial Setup section.

Change Alarm Ventilator Setting

Modify the Low Tidal Volume alarm setting to match the one shown below.

SETTING	VALUE
Low Vte/Low Vti	500 ml

Verify the Alarm

Wait up to 40 seconds and verify the following alarm signals:

- The High Priority audible indicator sounds
- A red light flashes on the Alarm Indicator/Audio Pause button
- The Low Tidal Volume alarm condition appears on the screen, highlighted in red

Modify Ventilator Alarm Settings

Modify the Low Tidal Volume alarm setting to match the one shown below.

SETTING	VALUE
Low Vte/Low Vti	50 ml

Verify Reset

Wait 40 seconds and verify the following auto-reset conditions:

- The High Priority audible indicator has stopped sounding
- The red light on the Alarm Indicator/Audio Pause button has stopped flashing

Restore Ventilator Settings

Modify the ventilator settings and change the following value shown below.

SETTING	VALUE
Low Vte/Low Vti	Off



VERIFY CIRCUIT DISCONNECT ALARM

This procedure verifies that the Circuit Disconnect alarm is working properly. It assumes that you have attached the test lung, verified ventilator settings, and turned on ventilator power as described in the Initial Setup section.

NOTE

The Low Inspiratory or Low Expiratory Pressure Alarm may also be detected.

Change Circuit Disconnect Ventilator Setting

Modify the Circuit Disconnect ventilator setting to match the value shown below.

SETTING	VALUE
Circuit Disconnect	10 seconds

Disconnect Test Lung

Disconnect the test lung from the circuit.

Verify the Alarm

Wait approximately 10 seconds and verify the following alarm signals:

- The High Priority Audible Indicator sounds
- A red light flashes on the Alarm Indicator/Audio Pause button
- The Circuit Disconnect alarm condition appears on the screen, highlighted in red

Reconnect Test Lung

Reconnect the test lung to the circuit.

Verify Reset

Wait at least 40 seconds and verify the following auto-reset conditions:

- The High Priority audible indicator has stopped sounding
- The red light on the Alarm Indicator/Audio Pause button has stopped flashing

Restore Ventilator Settings

Modify the ventilator settings and change the following values shown below.

SETTING	VALUE
Circuit Disconnect	Off

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VERIFY THE HIGH INSPIRATORY PRESSURE ALARM

This procedure verifies that the High Inspiratory Pressure alarm is working properly. It assumes that you have attached the test lung, verified ventilator settings, and turned on ventilator power as described in the Initial Setup section.

NOTE

If this alarm is not reset within 3 occurrences, the alarm is elevated to High Priority, and the High Priority Indicators occur.



Change Ventilator Settings

Modify the ventilator settings and change the following values shown below.

SETTING	VALUE
Mode	CV
Tidal Volume	500 ml
Breath Rate	12 BPM
Inspiratory Time	1.0 seconds
Flow Pattern	Ramp
PEEP	4 pressure units
Sigh	Off
Circuit Disconnect	Off
Low Inspiratory Pressure	6 pressure units
High Inspiratory Pressure	10 pressure units
Apnea	Off
All other alarms	Off

Verify the Alarm

Wait up to 40 seconds and verify the following alarm signals:

- The Medium Priority audible indicator sounds
- A yellow light flashes on the Alarm Indicator/Audio Pause button
- The High Inspiratory Pressure alarm condition appears on the screen, highlighted in yellow

Modify Ventilator Alarm Settings

Modify the High Inspiratory Pressure setting to match the one shown below.

SETTING	VALUE
High Inspiratory Pressure	60 pressure units

Verify Reset

Wait 40 seconds and verify the following auto-reset conditions:

- The Medium Priority audible indicator has stopped sounding
- The yellow light on the Alarm Indicator/Audio Pause button has stopped flashing

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VERIFY THE LOW INSPIRATORY PRESSURE ALARM

This procedure verifies that the Low Inspiratory Pressure alarm is working properly. It assumes that you have attached the test lung, verified ventilator settings, and turned on ventilator power as described in the Initial Setup section.

Change Ventilator Settings

Modify the ventilator settings and change the following values shown below.

SETTING	VALUE
Mode	CV
Tidal Volume	500 ml
Breath Rate	12 BPM
Inspiratory Time	1.0 seconds
Flow Pattern	Ramp
PEEP	4 pressure units
Sigh	Off
Circuit Disconnect	Off
Low Inspiratory Pressure	40 pressure units
High Inspiratory Pressure	60 pressure units
Apnea	Off
All other alarms	Off

Verify the Alarm

Wait up to 40 seconds and verify the following alarm signals:

- The High Priority audible indicator sounds
- A red light flashes on the Alarm Indicator/Audio Pause button
- The Low Inspiratory Pressure alarm condition appears on the screen, highlighted in red

Modify Ventilator Alarm Settings

Modify the Low Inspiratory Pressure setting to match the one shown below.

SETTING	VALUE
Low Inspiratory Pressure	6 pressure units

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Verify Reset

Wait 40 seconds and verify the following auto-reset conditions:

- The High Priority audible indicator has stopped sounding
- The red light on the Alarm Indicator/Audio Pause button has stopped flashing

10.3.5 BATTERY FUNCTION VERIFICATION

Make sure the batteries are functioning properly and fully charged before patient use.

VERIFY THE DETACHABLE AND INTERNAL (LITHIUM-ION) BATTERIES FUNCTION

- 1. Connect AC Power to the device and verify that the green AC LED on the front panel is lit.
- 2. Verify that the detachable battery is properly installed.
- 3. Turn the device on and verify that both the detachable and internal battery symbols appear on the display. Verify that if either battery is less than fully charged, the charge symbol will display on the respective battery.
- 4. Disconnect the AC Power source from the device.
 - Verify that the AC Power Disconnected alarm message appears on the display and the green AC LED is not lit. Press Reset.
 - Verify that the detachable battery symbol shows the level of charge noted in the previous step and that the device continues to operate.
 - Verify that the detachable battery symbol has a black box around it to indicate that it is in use.
- 5. Disconnect the detachable battery pack from the device.
 - Verify that the Detach Batt Disconnected alarm message appears on the display. Press Reset.
 - Verify that the internal battery symbol shows the same level of charge as noted in Step C and the device continues to operate.
 - Verify that the internal battery symbol has a black box around it to indicate that it is in use.
- 6. Reconnect the Detachable Battery and AC Power source.

VERIFY THE EXTERNAL BATTERY FUNCTION (IF AVAILABLE)

- 1. Connect AC Power to the device and verify that the green AC LED is lit.
- 2. Connect the external battery cable to the external battery and to the ventilator.
- 3. Verify that the external battery symbol is shown on the display and some level of charge is present.
- 4. Disconnect the AC Power source from the device.
 - Verify that the AC Power Disconnected alarm message appears on the display and the green AC LED is not lit. Press Reset.
 - Verify that the external battery symbol shows the level of charge as noted in the previous step and the device continues to operate.
 - Verify that the external battery symbol has a black box around it to indicate that it is in use.
- 5. Reconnect the AC Power source.

ALARM AND EVENT LOG CLEAN-UP

- In the Setup Menu, select Alarm Log.
 - a. Press Clear to clear the log file.
 - b. Press Yes to confirm.



- c. Press Finish to complete.
- 2. In the Setup Menu, select Event Log.
 - a. A. Press Clear to clear the log file.
 - b. Press Yes to confirm.
 - c. Press Finish to complete.

RESULTS

All portions of this checkout procedure should be completed prior to connection to the patient. If any of the tests fail to complete as indicated, if possible, correct the error, clear the alarm and resume testing. If correction of the failed portion is not possible, return the device to Respironics or an authorized service center for service and repair.

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10.4 TRILOGY 200 CHECKOUT PROCEDURE DATA SHEET

10.4.1 VISUAL INSPECTION

	YES	NO
Damaged Parts?		

10.4.2 SETTING & ALARM TESTS

High Tidal Volume Alarm Setting?	PASS	FAIL
Low Tidal Volume Alarm Setting?	PASS	FAIL
Circuit Disconnect Alarm Setting?	PASS	FAIL
High Inspiratory Pressure Alarm Setting?	PASS	FAIL
Low Inspiratory Pressure Alarm Setting?	PASS	FAIL
Battery Function Verification?	PASS	FAIL

Signature:	 	
Date:	 	
Sorial Number		



10.5 TRILOGY O2 & TRILOGY 202 CHECKOUT PROCEDURE

This chapter details the test procedures that should be performed by the clinician prior to connecting the device to the patient. Test the Active PAP, Active Flow, and Passive circuit types if you want to do a complete checkout on the device. The tests should be performed as described in order to verify proper operation of the device. Some of the procedures in this chapter require you to change settings on the device.

10.5.1 Tools Required

- Universal Porting Block
- Active PAP Exhalation Device
- Active Flow Exhalation Device with Flow Sensor
- Trilogy Universal Active PAP Tube Adapter
- Whisper Swivel II
- Test Lung
- High Pressure O₂ Hose
- Oxygen Monitor
- Small Flat Head Screwdriver

10.5.2 VISUAL INSPECTION

- 1. Verify that the enclosure is not broken and that all applicable screws are in place.
- 2. Verify that the device handle, SD Card door, and detachable battery are secure and in good working order.
- 3. Verify that the rubber feet are on the bottom of the device.

10.5.3 INITIAL SETUP

- 1. Connect the power cord to the device and then to an AC outlet.
- 2. Attach the test lung to the patient connection end of the desired circuit (Active PAP, Active Flow, or Passive).
- 3. Connect the device to a suitable high pressure O₂ source.
- 4. Connect and set up an external O₂ monitor per the manufacturer's instruction manual.
- 5. Follow the instructions in Chapter 5 to access the Setup Screen.



10.5.4 SETTINGS AND ALARMS TESTS

Complete the following steps to set up the settings and alarms tests.

SETUP

Settings And Alarms Menu

Modify the settings in the Settings and Alarms menu to match those shown below.

SETTING	VALUE
Dual Prescription	Off
Circuit Type	Active PAP, Active Flow, or Passive
Therapy Mode	S/T
AVAPS (passive circuit only)	Off
IPAP	20 pressure units
EPAP	4 pressure units
FiO ₂	45%
Breath Rate	12 BPM
Inspiratory Time	1.6 seconds
Trigger Type (passive circuit type)	Auto-Trak
Flow Trigger Sensitivity (Active PAP or Active Flow circuit type)	6.0 l/min
Leak Compensation (Active Flow circuit type)	On
Flow Cycle Sensitivity (Active PAP or Active Flow circuit type)	20%
Rise Time	1
Ramp Length	Off
All other alarms	Off



Options Menu

Modify the settings in the Options menu to match those shown below.

SETTING	VALUE
Menu Access	Full
Detailed View	On
All other settings	Discretionary

Turn Device Power On

Press the Start/Stop button on the front of the ventilator. The system will begin operating using the defined ventilation settings.

VERIFY THE HIGH TIDAL VOLUME ALARM

This procedure verifies that the High Tidal Volume alarm is working properly. For Passive and Active Flow circuits, this will verify the High Vte alarm. For Active PAP circuits, this will verify the High Vti alarm. It assumes that you have attached the test lung, verified the ventilator settings, and turned on ventilator power as described in the Initial Setup section.

NOTE

Do not use the "reset" button to manually reset the alarm. Instead, use the "Modify" button to change ventilator settings. This note applies to all tests.

Change Alarm Ventilator Setting

Modify the High Tidal Volume alarm setting to match the one shown below.

SETTING	VALUE
High Vte/High Vti	50 ml

Verify the Alarm

Wait up to 40 seconds and verify the following alarm signals:

- · The High Priority audible indicator sounds
- A red light flashes on the Alarm Indicator/Audio Pause button
- The High Tidal Volume alarm condition appears on the screen, highlighted in red

Modify Ventilator Alarm Settings

Modify the High Tidal Volume alarm setting to match the one shown below.

SETTING	VALUE
High Vte/High Vti	500 ml



Verify Reset

Wait 40 seconds and verify the following auto-reset conditions:

- The High Priority audible indicator has stopped sounding
- The red light on the Alarm Indicator/Audio Pause button has stopped flashing

Restore Ventilator Settings

Modify the ventilator settings and change the following value shown below.

SETTING	VALUE
High Vte/High Vti	Off

VERIFY THE LOW TIDAL VOLUME ALARM

This procedure verifies that the Low Tidal Volume alarm is working properly. For Passive and Active Flow circuits, this will verify the Low Vte alarm. For Active PAP circuits, this will verify the Low Vti alarm. It assumes that you have attached the test lung, verified the ventilator settings, and turned on ventilator power as described in the Initial Setup section.

Change Alarm Ventilator Setting

Modify the Low Tidal Volume alarm setting to match the one shown below.

SETTING	VALUE
Low Vte/Low Vti	500 ml

Verify the Alarm

Wait up to 40 seconds and verify the following alarm signals:

- The High Priority audible indicator sounds
- A red light flashes on the Alarm Indicator/Audio Pause button
- The Low Tidal Volume alarm condition appears on the screen, highlighted in red

Modify Ventilator Alarm Settings

Modify the Low Tidal Volume alarm setting to match the one shown below.

SETTING	VALUE
Low Vte/Low Vti	50 ml

Verify Reset

Wait 40 seconds and verify the following auto-reset conditions:

- The High Priority audible indicator has stopped sounding
- The red light on the Alarm Indicator/Audio Pause button has stopped flashing

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Restore Ventilator Settings

Modify the ventilator settings and change the following value shown below.

SETTING	VALUE
Low Vte/Low Vti	Off

VERIFY CIRCUIT DISCONNECT ALARM

This procedure verifies that the Circuit Disconnect alarm is working properly. It assumes that you have attached the test lung, verified ventilator settings, and turned on ventilator power as described in the Initial Setup section.

NOTE
The Low Inspiratory or Low Expiratory Pressure Alarm may also be detected.

Change Circuit Disconnect Ventilator Setting

Modify the Circuit Disconnect ventilator setting to match the value shown below.

SETTING	VALUE
Circuit Disconnect	10 seconds

Disconnect Test Lung

Disconnect the test lung from the circuit.

Verify the Alarm

Wait approximately 10 seconds and verify the following alarm signals:

- The High Priority Audible Indicator sounds
- A red light flashes on the Alarm Indicator/Audio Pause button
- The Circuit Disconnect alarm condition appears on the screen, highlighted in red

Reconnect Test Lung

Reconnect the test lung to the circuit.

Verify Reset

Wait at least 40 seconds and verify the following auto-reset conditions:

- The High Priority audible indicator has stopped sounding
- The red light on the Alarm Indicator/Audio Pause button has stopped flashing



Restore Ventilator Settings

Modify the ventilator settings and change the following values shown below.

SETTING	VALUE
Circuit Disconnect	Off

VERIFY THE HIGH INSPIRATORY PRESSURE ALARM

This procedure verifies that the High Inspiratory Pressure alarm is working properly. It assumes that you have attached the test lung, verified ventilator settings, and turned on ventilator power as described in the Initial Setup section.

NOTE

If this alarm is not reset within 3 occurrences, the alarm is elevated to High Priority, and the High Priority Indicators occur.

Change Ventilator Settings

Modify the ventilator settings and change the following values shown below.

SETTING	VALUE
Mode	CV
Tidal Volume	500 ml
FiO ₂	21%
Breath Rate	12 BPM
Inspiratory Time	1.0 seconds
Flow Pattern	Ramp
PEEP	4 pressure units
Sigh	Off
Circuit Disconnect	Off
Low Inspiratory Pressure	6 pressure units
High Inspiratory Time	10 pressure units
Apnea	Off
All other alarms	Off

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Verify the Alarm

Wait up to 40 seconds and verify the following alarm signals:

- The Medium Priority audible indicator sounds
- A yellow light flashes on the Alarm Indicator/Audio Pause button
- The High Inspiratory Pressure alarm condition appears on the screen, highlighted in yellow

Modify Ventilator Alarm Settings

Modify the High Inspiratory Pressure setting to match the one shown below.

SETTING	VALUE
High Inspiratory Pressure	60 pressure units

Verify Reset

Wait 40 seconds and verify the following auto-reset conditions:

- · The Medium Priority audible indicator has stopped sounding
- The yellow light on the Alarm Indicator/Audio Pause button has stopped flashing

VERIFY THE LOW INSPIRATORY PRESSURE ALARM

This procedure verifies that the Low Inspiratory Pressure alarm is working properly. It assumes that you have attached the test lung, verified ventilator settings, and turned on ventilator power as described in the Initial Setup section.



Change Ventilator Settings

Modify the ventilator settings and change the following values shown below.

SETTING	VALUE
Mode	CV
Tidal Volume	500 ml
FiO ₂	21%
Breath Rate	12 BPM
Inspiratory Time	1.0 seconds
Flow Pattern	Ramp
PEEP	4 pressure units
Sigh	Off
Circuit Disconnect	Off
Low Inspiratory Pressure	40 pressure units
High Inspiratory Time	60 pressure units
Apnea	Off
All other alarms	Off

Verify the Alarm

Wait up to 40 seconds and verify the following alarm signals:

- The High Priority audible indicator sounds
- A red light flashes on the Alarm Indicator/Audio Pause button
- The Low Inspiratory Pressure alarm condition appears on the screen, highlighted in red

Modify Ventilator Alarm Settings

Modify the Low Inspiratory Pressure setting to match the one shown below.

SETTING	VALUE
Low Inspiratory Pressure	6 pressure units

Verify Reset

Wait 40 seconds and verify the following auto-reset conditions:

- The High Priority audible indicator has stopped sounding
- The red light on the Alarm Indicator/Audio Pause button has stopped flashing

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VERIFY THE O₂ BLENDING OPERATION

This procedure verifies that the Low Oxygen Flow alarm and the Low Oxygen Inlet Pressure alarm are working properly. These alarms apply to all circuit types. It assumes that you have attached the test lung, verify the ventilator settings, and turned on the ventilator power as described in the Initial Setup section.

Setup

- 1. Connect an external O₂ Monitor (in accordance with the recommended manufacture's guidelines) in-line with the patient tube. Make sure the O₂ monitor is properly calibrated before proceeding.
- 2. Connect the ventilator's oxygen input port to a source of high pressure O₂ (60 psi nominal). Turn on O₂ flow to the ventilator.

Ventilator Settings

Set the ventilator FiO₂ setting to 45%.

Verify Blending

- 1. Turn on the ventilator.
- 2. Verify the set level of FiO₂ is satisfied using an external O₂ monitor.

Verify alarm

- 1. Shut off or disconnect the source of high pressure O₂ to the ventilator.
- 2. Wait 1 minute and verify the following alarm signals:
- · The Priority audible indicator sounds.
- A red light flashes on the Alarm Indicator/Audio Pause Button.
- The following alarm conditions appear on the screen, highlighted in red:
 - Low Oxygen Flow
 - Low Oxygen Inlet Pressure



Verify Reset

- 1. Reconnect or turn on the source of high pressure O_2 to the ventilator.
- 2. Wait 1 minute and verify the following:
- The set level of FiO₂ is satisfied using an external O₂ monitor.
- The High Priority audible indicator has stopped sounding.
- The red light on the Alarm Indicator/Audio Pause button has stopped flashing.

10.5.5 BATTERY FUNCTION VERIFICATION

Make sure the batteries are functioning properly and fully charged before patient use.

VERIFY THE DETACHABLE AND INTERNAL (LITHIUM-ION) BATTERIES FUNCTION

- 1. Connect AC Power to the device and verify that the green AC LED on the front panel is lit.
- 2. Verify that the detachable battery is properly installed.
- Turn the device on and verify that both the detachable and internal battery symbols appear on the display. Verify that if either battery is less than fully charged, the charge symbol will display on the respective battery.
- 4. Disconnect the AC Power source from the device.
 - Verify that the AC Power Disconnected alarm message appears on the display and the green AC LED is not lit. Press Reset.
 - Verify that the detachable battery symbol shows the level of charge noted in the previous step and that the device continues to operate.
 - Verify that the detachable battery symbol has a black box around it to indicate that it is in use.
- 5. Disconnect the detachable battery pack from the device.
 - Verify that the Detach Batt Disconnected alarm message appears on the display. Press Reset.
 - Verify that the internal battery symbol shows the same level of charge as noted in Step C and the device continues to operate.
 - Verify that the internal battery symbol has a black box around it to indicate that it is in use.
- 6. Reconnect the Detachable Battery and AC Power source.

VERIFY THE EXTERNAL BATTERY FUNCTION (IF AVAILABLE)

- 1. Connect AC Power to the device and verify that the green AC LED is lit.
- 2. Connect the external battery cable to the external battery and to the ventilator.
- 3. Verify that the external battery symbol is shown on the display and some level of charge is present.
- 4. Disconnect the AC Power source from the device.
 - Verify that the AC Power Disconnected alarm message appears on the display and the green AC LED is not lit. Press Reset.
 - Verify that the external battery symbol shows the level of charge as noted in the previous step and the device continues to operate.
 - Verify that the external battery symbol has a black box around it to indicate that it is in use.
- 5. Reconnect the AC Power source.

ALARM AND EVENT LOG CLEAN-UP

In the Setup Menu, select Alarm Log.



- a. Press Clear to clear the log file.
- b. Press Yes to confirm.
- c. Press Finish to complete.
- 2. In the Setup Menu, select Event Log.
 - a. A. Press Clear to clear the log file.
 - b. B. Press Yes to confirm.
 - c. Press Finish to complete.

RESULTS

All portions of this checkout procedure should be completed prior to connection to the patient. If any of the tests fail to complete as indicated, if possible, correct the error, clear the alarm and resume testing. If correction of the failed portion is not possible, return the device to Respironics or an authorized service center for service and repair.

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10.6 TRILOGY O_2 & TRILOGY 202 CHECKOUT PROCEDURE DATA SHEET

10.6.1 VISUAL INSPECTION

	YES	NO
Damaged Parts?		

10.6.2 SETTING & ALARM TESTS

High Tidal Volume Alarm Setting?	PASS	FAIL
Low Tidal Volume Alarm Setting?	PASS	FAIL
Circuit Disconnect Alarm Setting?	PASS	FAIL
Adm octang.		
High Inspiratory	PASS	FAIL
Pressure Alarm Setting?		
Low Inspiratory	PASS	FAIL
Pressure Alarm Setting?		
	•	
Verify the O ₂	PASS	FAIL
Blending Operation?		
Battery Function?	PASS	FAIL

Signature:	 	 	
Date:	 	 	
Serial Number:			



10.7 DOWNLOADING THE TRILOGY FIELD SERVICE APPLICATION, TRILOGY TOOLBOX SOFTWARE, AND NATIONAL INSTRUMENTS DAQMX SOFTWARE

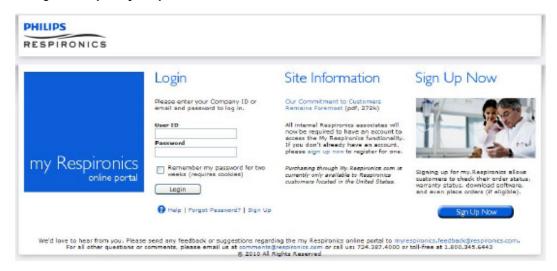
NOTE

Respironics service software is now available at http://my.respironics.com. In the event that you are unable to access this site, log onto http://servicesoftware.respironics.com. to download Respironics service software.

You must be a registered user and have completed the Trilogy Certified Training Course in order to download the *Trilogy Field Service Application, Trilogy Toolbox Software, and National Instruments DAQmx Software.* To become a registered user and to download Trilogy Software and Documentation, you must successfully complete the Trilogy Service Training class.

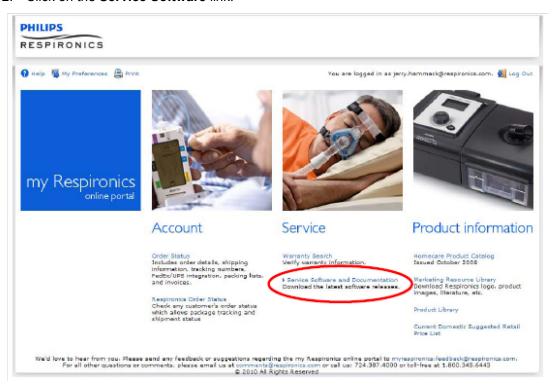
Once you have access to download the software, perform the following:

1. Log into http://my.respironics.com.





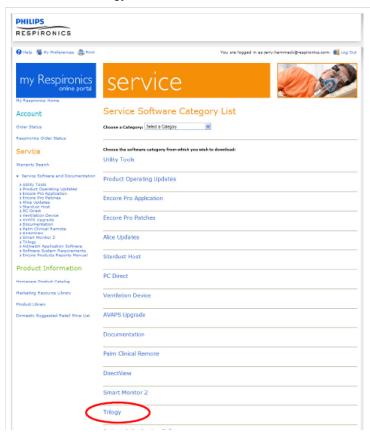
2. Click on the Service Software link.



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3. In the menu below, select Trilogy.



4. Click on the **Download** button adjacent to the software you wish to download - *Trilogy Toolbox, Trilogy Field Service Application Software, or National Instruments DAQmx.*

NOTE

Remember to periodically log onto http://my.respironics.com and check for Trilogy software and firmware updates.

IMPORTANT!

The programs may be downloaded in any order, but the applications must be installed to the computer in the order of FSA then TBA then DAQ

IMPORTANT!

Install the NI DAQmx software after the Field Service Application and the Trilogy Toolbox have been installed on your PC. When installing the NI DAQmx software you will see a series of onscreen prompts. Use all of the default settings associated with the on-screen prompts until you get to the software version screen. For NI DAQmx to interface properly with the Trilogy Field Service Application and Trilogy Toolbox, Version 7.1 MUST be selected.



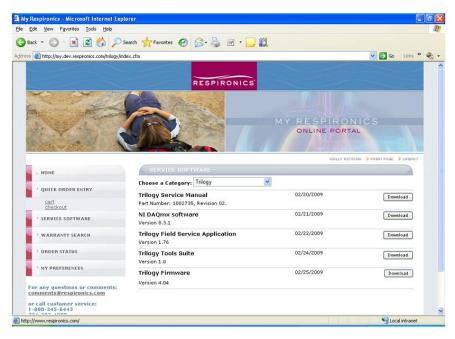


FIGURE 10-1: DOWNLOADING SOFTWARE SAMPLE SCREEN

IMPORTANT NOTE

If the Trilogy Toolbox is already installed on your PC, clicking on the download button will remove the current version of the software from the PC. Verify that the installed version is the latest version, otherwise, you must perform the "Download" procedure twice.

At anytime that you update any of the software applications, all applications must be completely removed using the ADD/REMOVE PROGRAMS feature of Windows; then all applications must be installed in the following order: FSA, TBA, and DAQ-mx.

When you click on the **Download** button, the "Run or Save?" window will appear.

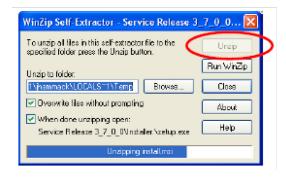
- 5. Click on **Save** to download the software and save it to a specific location on your PC.
- 6. Follow the on-screen prompts to "Save" the software.
- 7. Once the downloads are completed, next begin the installation process for each of the applications. Remember that the applications must be installed to the computer hard drive in the proper order (FSA, TBA, DAQ) or errors will occur when trying to run the installed applications.
- 8. Open the folder that the applications were downloaded to.
- 9. Select the "Service Release_x_x_x_x.exe" file first, where the x's represent the current version of FSA, by double clicking with the mouse on the application.

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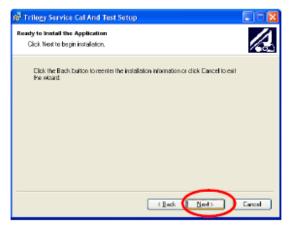
10. Follow the prompts and select the options highlighted by the RED Circles.

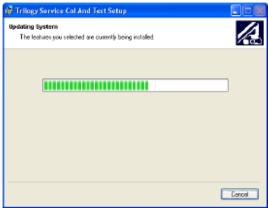






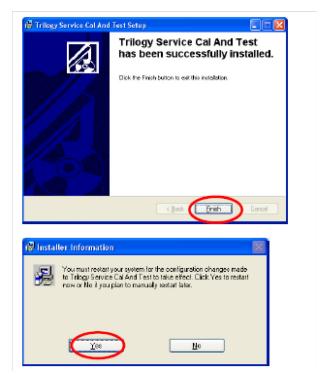






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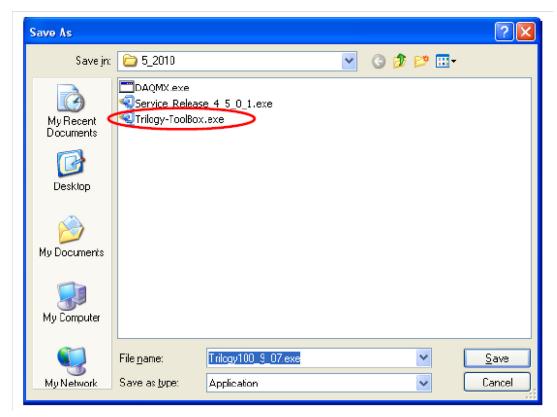
11. You must restart your computer after this installation. Once the computer restarts there will be 4 new loons on the Desktop screen of the computer.



- 12. Two portions of the Trilogy testing, a Pre Test which provides calibration and verification prior to the Run-in procedure and the Post Test portion which provides verification after the Run-in procedure and during the Preventative Maintenance procedure.
- 13. There are also two ways to perform the testing, with Manual Flow testing you must use Shop Air, Compressed (Cylinder) Air, or Compressed (Cylinder) O2.



14. Next from the folder, select the TBA for installation by double clicking with the mouse the "Trilogy-ToolBox.exe" file.



15. Follow the prompts and select the options highlighted by the RED Circles.

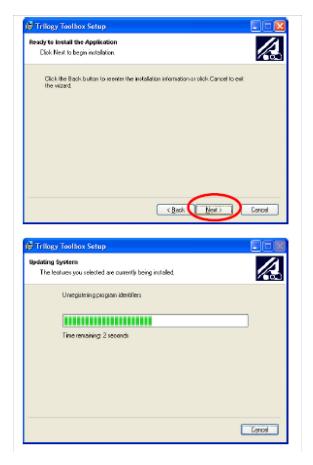


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16. At this screen, the installation will ask for the folder to install the LogDogg Application into. The Default folder needs to be modified by inserting the additional folder in the install path.



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17. Follow the Red circle examples below.



18. Change from the default listed above to the new path listed below.

