



Clearing The Airway Is Our #1 Priority

Operating Instructions & Maintenance Manual

SSCOR[®] DUET[®]
Model 2314 Series



SSCOR INC.

2125 N. Madera Rd. Unit C
Simi Valley, CA 93065 USA

www.sscor.com
shop.sscor.com
Email: marketing@sscor.com
techsupport@sscor.com

Tel 1+818-504-4054



Intertek



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Caution Notice

1. SSCOR suction units are not designed or intended for use in extended procedures that require prolonged high vacuum/low airflow applications, as is the case in wound drainage or endoscopic use or in any other procedure that produces high vacuum levels within an occluded system for an extended period of time. Turn the suction unit off when it is not in use.
2. Federal law restricts this device to sale, distribution, and use by, or on the order of a physician, emergency medical technician, or other medical practitioner. For use by medical personnel trained in suctioning techniques and in the use of medical suction equipment.
3. Operator should be thoroughly familiar with these operating instructions before this device is used.
4. Do not use in the presence of flammable agents or anesthetics.
5. Before testing for vacuum over -300mmHg look for an expiration date on the canister (where applicable) and change the canister if the canister has passed the expiration date to minimize the possibility of implosion, which can occur when a canister is aged or damaged.
6. External equipment intended for connection to signal input, signal output or other connectors, shall comply with relevant IEC standard (e.g. IEC 60950 for IT equipment and the IEC 60601 series for medical electrical equipment). In addition, all such combinations–systems – shall comply with the standard IEC 60601-1-1, safety requirements for medical electrical systems. Equipment not complying with IEC 60601 shall be kept outside the patient environment, as defined in the standard. Any person who connects external equipment to signal input, signal output or other connectors has formed a system and is therefore responsible for the system to comply with the requirements of IEC 60601-1-1. If in doubt, contact qualified technician or your local representative.
7. A blinking LED on the control panel indicates low battery capacity. This means the battery has been subjected to irreparable damage. REPLACE THE BATTERY! For the best quality, we recommend purchasing battery replacements from SSCOR (Available to purchase at shop.sscor.com, by sending an email to sales@sscor.com, or by calling +1 818-504-4054).
8. We strongly recommend original SSCOR parts and recommend against using third-party components including batteries.
9. Where the integrity of the external protective earth conductor arrangement (ground) is in doubt, the unit shall be operated from its internal electrical power source (battery only). Grounding reliability can only be achieved when connected to an equivalent receptacle marked "Hospital Only" or "Hospital Grade".
10. Dispose of device according to local / regional / national requirements for the disposal of electronic waste at the end of the expected service life.
11. Dispose of single use accessories according to local / regional / national requirements for the disposal of hazardous waste.
12. When running the device from the AC power supply do not position the device in a way that makes it difficult to unplug the power cord from the device.
13. SSCOR aspirators are intended for use in various electromagnetic environments typified by hospital and EMS situations and public establishments that use standard mains power. It is recommended that the user test SSCOR aspirators in conjunction with other devices that may be in operation at the time of use. SSCOR aspirators have very low RF emissions and are not likely to cause interference in nearby electronic equipment. SSCOR also recommends the other devices in the environment at the time of use also meet IEC 60601EMC/RF requirements.

SSCOR Duet®

Model 2314 Series - AC suction unit with battery back-up

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General Description

SSCOR suction units are designed for hospital crash carts, patient transport and emergency medical service. Battery condition is automatically monitored and visualized by indicator lights on the control panel. The battery should be charged to a dependable working charge in 6 to 8 hours when connected to the charging source. Suction power can be regulated when full power may be considered harmful to the patient. SSCOR suction units are designed to provide instant, effective suctioning, independent of external sources of power and can be pre-set to be activated immediately upon reaching the distressed patient. All controls are clearly labeled and easily accessible.

The SSCOR Duet is a portable, 100V-240V AC constant suction device, with a 12V DC battery back-up. The battery is charged by an internal, DC dual mode battery charger. The charger monitors the battery, charges the battery only when necessary, shuts down the unit if the battery is low and signals battery condition. A fully charged battery at full capacity will power the unit for 45 minutes ($\pm 10\%$).



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Standard Canister Options

The SSCOR Duet is available in two standard canister configurations, the 2314 and the 2314H, to specifically fit the canister you use in your facility.



Duet with 1200cc Bemis Canister - pn:2314 (Canister is included)



Duet with 1200cc Medline Canister - pn:2314H (Canister is not included)



Duet with 1200cc Cardinal Canister pn:2314H (Canister is not included)



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Warranty

SSCOR warrants that each new product is free from defects in material and workmanship under normal use and service for a period of five years from date of original purchase from SSCOR, Inc. If returned to SSCOR, we will arrange for repairs or replacement within the terms of the warranty. The product should be decontaminated and returned properly packaged and postage prepaid. Loss or damage in transit to the factory shall be at the purchaser's risk. Loss or damage of return shipment from SSCOR shall be at the purchaser's risk. Please go to the support page at www.SSCOR.com to request a return authorization number.

Conditions:

This warranty covers none of the following:

1. The warranty shall not apply to any SSCOR device in which a component has been altered; determined to be using a non SSCOR component; or altered in any way so as, in SSCOR's judgment, to affect its safety or efficacy.
2. SSCOR product which has been subject to misuse, negligence, or accident.
3. The serial number has been altered, effaced or removed.
4. SSCOR product which has been setup otherwise than in accordance with the instructions furnished by SSCOR.
5. Maintenance and repair due to normal wear and tear.
6. Installation of product in a way not shown in instruction for use furnished by SSCOR.
7. Faults in the system which SSCOR product is installed.
8. Batteries, Disposables items (canisters, patient tubing, and catheters), replacement parts, cosmetic irregularities that do not degrade the safety and efficacy of the SSCOR product.

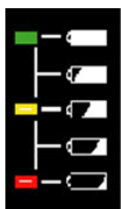
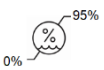
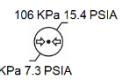
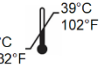
This warranty is in lieu of all other warranties expressed or implied and of all other obligations or liabilities on SSCOR's part, and SSCOR neither assumes, nor authorizes any representative or other persons to assume for it, any other liability in connection with the sale of SSCOR products.

This warranty gives you specific legal rights and you may also have other rights that vary from jurisdiction to jurisdiction. For countries where minimum warranty terms are determined by statute, the warranty term is the longer of the statutory period or the term listed above

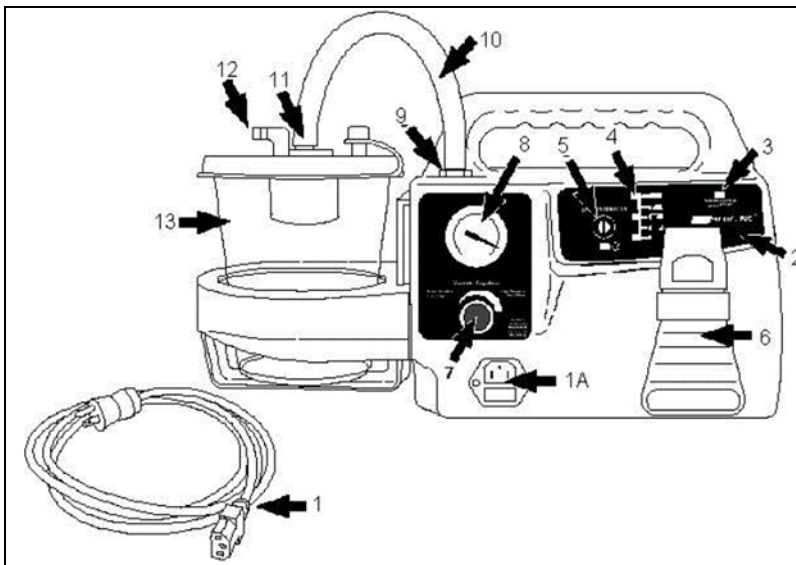
Model 2314 Series Battery Replacement Policy

SSCOR will replace any SSCOR Duet battery which fails to operate the pump to specifications for a period of one year from the date of purchase. Verify the condition of the battery per the battery test on Page 8 in this Operations Manual.

Description of Symbols

SYMBOL	LOCATION	MEANING
	Control Panel	Battery Level of Charge Green LED = Full Green / Yellow LED = High Yellow LED = Half Yellow / Red LED = Low Red LED = Battery Depleted Put device back on charge!
	Control Panel	Push On / Push Off
	Serialized Label	Attention—Consult Accompanying Documents
	Serialized Label	Separate collection for electronic equipment
	Serialized Label	Type BF Equipment
	Shipping Carton	Manufacturer
	Shipping Carton	Date of Manufacture
	Shipping Carton	Transport Storage Range
	Shipping Carton	Humidity
	Shipping Carton	Atmospheric Pressure
	Serialized Label	Operating Temperature Range
IP34	Serialized Label	The degree of protection provided by the chassis according to IEC 60529
	Regulator Label	Suction
	Control Panel	Pump is On
	Control Panel	Power Source Connected
100-240V ~	Serialized Label	Alternating Current (100-240V AC)

Getting Acquainted



#	DESCRIPTION	SKU
#1	Power Cord	80551-100
#1A	AC Wiring Harness	80971
#2	Control Panel	80023-115
#3	"Power Source Connected" LED	
#4	Battery Charge Indicator LEDs	
#5	ON/OFF Switch	
#6	Tubing Strap	50176
#7	Vacuum Regulator	22012-
#8	Vacuum Gauge	KIT2314
#9	Union Barb	90157
#10	Connector Tubing	50159-9
#11	Canister Vacuum Port	N/A
#12	Elbow Barb	42060
#13	1200cc Suction Canister	44241

Running the unit and Charging the Internal Battery from AC Power

1. Connect the female plug on the AC power cord **#1** to the AC receptacle on the device **#1A**.
2. Connect the male plug on the AC power cord **#1** to a grounded wall receptacle.
3. Check the control panel **#2**. The "power source connected" orange LED **#3** indicates a good connection to the AC power source and indicates the battery is charging. The battery condition indicators **#4** show the charge level of the battery (See Description of Symbols page 4).
4. Press the ON/OFF switch **#5** on the control panel to start the vacuum pump.

Running the unit from DC power

1. Disconnect the power cord **#1** and press the ON/OFF switch **#5** on the control panel to start the vacuum pump.
2. Check battery condition indicators **#4**

Adjusting the vacuum level

1. Occlude the patient tube and turn the vacuum regulator **#7** clockwise to the stop.
2. Observe gauge **#8** (located above the vacuum regulator). The vacuum reading should rise to -300mmHg from zero in 3 seconds. It should max out at approximately $\geq -525\text{mmHg}$. Lower levels of negative pressure will be observed at altitude.
3. Adjust the vacuum reading to your desired level. If the vacuum does not meet or exceed -525mmHg, check for a leak in the system i.e. tubing **#10**, barb connections (**#9,#11,#12**) or loose canister lid. Refer internal vacuum problems to qualified personnel.
4. After using the device, always reconnect the unit to the charging source; check the power source connected LED **#3**.

Battery Test

Run the following test routinely to ensure proper performance of the device

1. Plug the device in to charge for at least 8 hours before testing. The battery condition LED on the control panel should light green when the device is plugged in and the battery is fully charged.
2. To begin the test, make sure the tubing from the device to the canister is connected to the canister (1) and the canister lid is secure (2). Unplug the device from the external power source and turn the device on.



3. Check for vacuum by occluding the patient tube and set the vacuum regulator (#7) to the maximum vacuum setting.
4. Observe the regulator gauge (#8). The vacuum reading should rise to -300mmHg from zero in less than 3 seconds. It should max out at $\geq -525\text{mmHg}$. Un-occlude the patient tubing.
5. Allow the unit to run for 15 minutes on DC power. If the unit stops or slows during the 15 minutes, or if the battery condition indicators (#4) begin to blink, it is possible the battery capacity has been depleted. It is time to replace the battery.
6. With the regulator still at maximum vacuum, repeat steps 3 and 4.
7. If the unit is still running at full power after 15 minutes, adjust the regulator to the desired setting, turn the device off and put it back on charge.

SSCOR recommends replacement of the battery after 3 years.

Replacement battery may be purchased at shop.sscor.com, by sending an email to sales@sscor.com, or by calling +1 818-504-4054)

Environmental Conditions

Operating Temperature Range:	0 °C (+32 °F) – +39 °C (+102 °F)
Operating Relative Humidity:	0 - 93% (non condensing)
Operating Atmospheric Pressure:	8.9 Psi (62 kPa) – 15.4 Psi (106 kPa)
Operating Altitude:	<2000m
Storage & Transport Temperature:	-40 °C (-40 °F) – +70 °C (+158 °F)
Storage & Transport Relative Humidity:	0 - 95% (non condensing)
Storage & Transport Atmospheric Pressure:	7.3 Psi (50 kPa) – 15.4 Psi (106 kPa)
Transient Operating Temperature Range:	-20 °C (-4 °F) – +50 °C (+122 °F)
Transient Humidity Range:	15% - 93% (non condensing)
Time to warm from minimum storage temperature to operating temperature: 30 minutes	
Time to cool from maximum storage temperature to operating temperature: 30 minutes	

Trouble Shooting



WARNING: DO NOT ATTEMPT TO SERVICE THIS EQUIPMENT IF YOU ARE NOT A QUALIFIED MEDICAL REPAIR TECHNICIAN

PROBLEM	POSSIBLE CAUSE	CORRECTIVE ACTION
Power Source Connected LED not lit	Fuse blown Internal electrical connections disconnected Power cord disconnected	Replace fuse in power supply or input receptacle Reconnect electrical connections Reconnect cord
Does not function when switch is turned on	Battery discharged Internal electrical connections disconnected	Reconnect to charging source Open unit and re-connect electrical connections
Blinking LED	Low battery capacity	Replace battery
No suction when pump is running	Vacuum line (10) loose Canister (13) defective Canister lid (13) loose Catheter thumb vent is open Debris has collected in the pump Defective Pump Defective regulator Tubing disconnected	Inspect pneumatic connections Replace canister Re-secure canister lid Occlude thumb vent on suction tip Replace pump Replace Regulator Reconnect tubing

General Specifications

CHARACTERISTICS	SPECIFICATIONS
Size	17"L x 9"H x 5.25"W(43.18cm L x 22.86cm H x 13.33cm W)
Weight	10.65 lbs (device only) 4.83 kgs (device only)
Vacuum Pump	12V DC. Clinical Airflow $\geq$ 30LPM Exceeds 525mmHg
Variable Regulator (7)	Controls negative pressure
Regulator Gauge (8)	Calibrated in mmHg. Color-coded
Power Source: Battery (DC Power) Power Supply (AC Power)	Rechargeable Sealed Lead Acid. SSCOR part # 80635 100V-240VAC, 47-63Hz Uses 1ea. 3A 250V fast acting fuse; Output 14.75 VDC
Switch (5)	Off/Standby/On membrane switch
AC receptacle (1A)	Uses (2) 2.5A 250V fast acting fuses
Collection Canister (13)	1200cc/ml SSCOR part #48041
Patient tubing	Vinyl tubing 9/32"ID x 72"L (7.1mm ID x 182.88 cm L) SSCOR part #43200
SSCOR Suction Catheter	SSCOR SDC Catheter SSCOR part # 200-00002 Or SSCOR HI-D Catheter SSCOR part # 44241

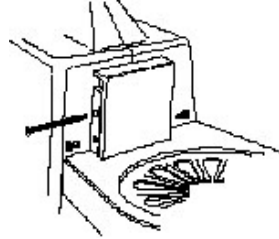


Internal Access



WARNING: Do not attempt to service this equipment if you are not a qualified medical repair technician. Take appropriate ESD abatement measures when handling all internal components.

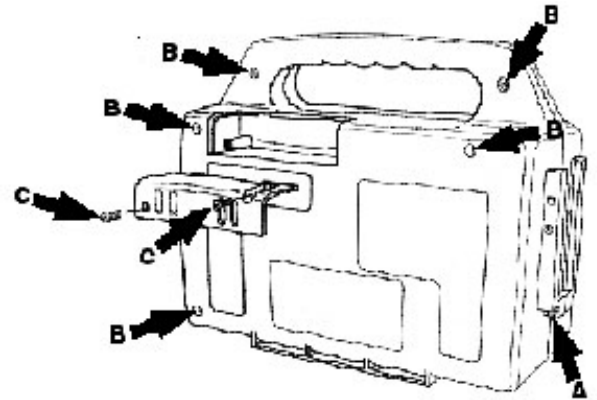
To open or close the unit, first remove the canister holder:
Remove the set screws on the canister bracket.
Lift the canister holder up and out of the main chassis.



The clamshell design enables access to internal components. All 6-32 Phillips head fastening screws are located on the back of the unit.

REMOVE MAIN CHASSIS SCREWS

- (A) 1 screw under the canister holder
- (B) 2 screws in the back of the handle
- (B) 3 screws in the back of the main chassis.
- (C) It should not be necessary to open the exhaust door unless fluids have entered the system



- (D) PC Board-Electrical Circuits (Internal Dual Mode Battery Charger).

Do not attempt to service the PC Board.

- (E) Power Supply. Do not attempt to service the Power Supply. Return to SSCOR for service.

- (F) Functional ground.

- (G) AC receptacle. Two 250V-2.5A fuses are in the receptacle fuse drawer.

- (H) Battery Bracket

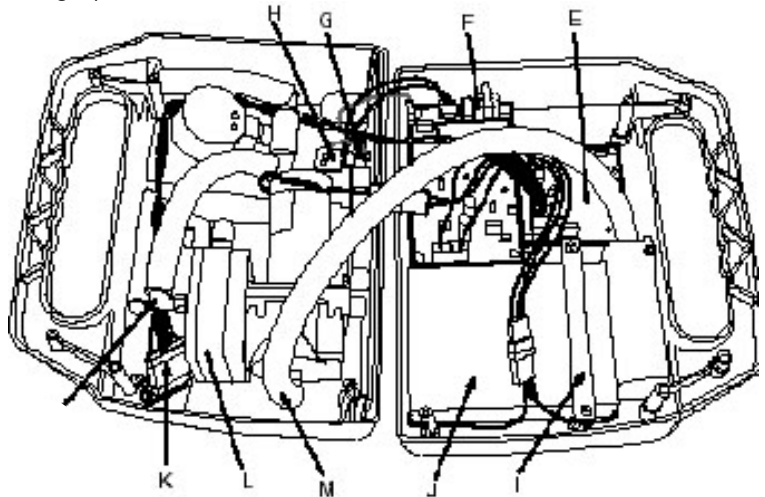
- (I) Battery, 12V, Sealed Lead Acid

- (J) Wiring connection to the control panel

- (K) Vacuum pump. Return to SSCOR for service. Do not attempt to service the vacuum pump.

- (L) Exhaust barb

- (M) Vacuum barb



Maintenance

Preventive Care

Observe the following maintenance routine to ensure readiness at any time:

1. When the SSCOR aspirator is not in use, keep batteries on continuous charge.
2. Test the SSCOR aspirator at regular intervals; See page 6.
3. Make sure the SSCOR aspirator is always clean and ready for use.
4. If the procedure produced an excessive quantity of fluids, check the vacuum line #10 for evidence of moisture. If the vacuum line between the pump and canister is moist, it is possible that fluids have reached the vacuum pump. See Disinfection Instructions below.

For technical assistance, replacement parts, and accessories, call +1 818-504-4054, email techsupportgrp@sscor.com, or visit shop.sscor.com.

Sanitation

As soon as possible after use, the single use disposable canister, patient tubing and catheter should be discarded according to local / regional / national requirements for the disposal of hazardous waste materials. Clean using a mild detergent and if necessary, disinfect with a mild disinfectant such as 10 to 1 bleach and rinse using clear water to remove any residue.

If the exhaust filter is soiled, remove the two screws holding the filter door in place (see page 8), remove the soiled filter and replace with new filter (SSCOR part number 90160).

Do not reuse any single use disposable parts; do not submerge the device into any liquid, this will void the warranty and cause the device to malfunction.

Disinfection

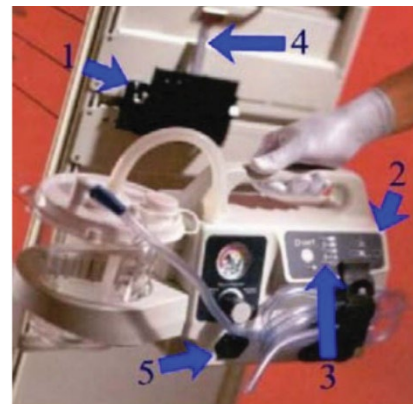
Use personal protective equipment such as gloves, a smock, and face and eye protection when handling units that are suspected to be contaminated.

Caution: Disconnect the unit from any power source prior to cleaning the unit. When cleaning the interior of the chassis, disconnect the battery from the PC Board to prevent damaging the PC Board.

Part	Cleaning and Disinfecting
Collection Canister	Disposable item, re-use not permitted. Use new canister for each patient.
Patient Tubing	Disposable item, re-use not permitted. Use new patient tubing for each patient.
SSCOR Suction Catheter	Disposable item, re-use not permitted. Use new SSCOR Suction catheter for each patient.
Vacuum Pump	Sterilization not permitted. Vacuum pump should be replaced if contaminated
Vacuum regulator	Sterilization not permitted. Vacuum pump should be replaced if contaminated
Electronic components	Sterilization not permitted. PC Board should be replaced if contaminated.
Plastic Chassis	Wipe with damp cloth or disinfectant wipe. Sterilization not permitted.

Bracket Mounting Instructions **SSCOR** Duet

If you purchased the Model 2314B and intend to mount the device to the side of a crash cart, mount the bracket to a suitable safe mounting surface using the four holes on the side of the bracket (1). Mount the bracket so the Duet suction unit can be released from the bracket easily and there is enough room to work the handle (4). The control panel (2) should be easily viewed and the battery condition LED's (3) visible when the unit (5) is connected to a grounded hospital receptacle. When placing the suction unit into the bracket, be sure the unit snaps into the bracket and is securely held in place.



Preventive Maintenance

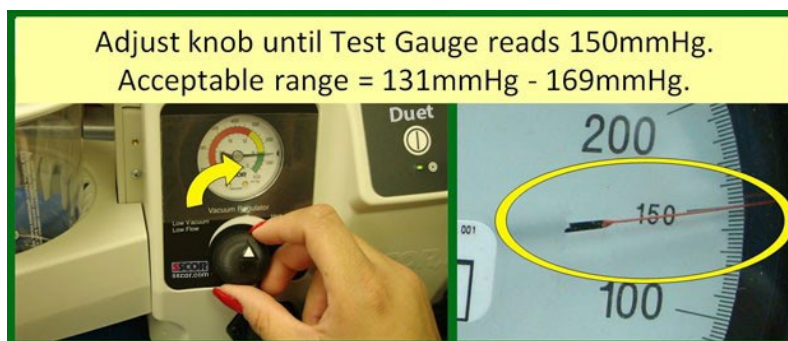
Perform the following preventive maintenance procedure at least annually.

1. When applicable, check the canister to determine if the expiration date has passed. If the expiration date has passed, replace the canister.
2. Test the health of the battery per the battery test on page 7:
For a further explanation of the **SSCOR** battery run test, visit the **SSCOR** Website at: http://www.sscor.com/training_videos.html.
3. Determine if the gauge is properly calibrated.



After the battery has been fully charged (when the battery condition LED on the control panel is green), connect tubing from the patient port of the canister to a calibrated negative pressure test gauge. Adjust the vacuum pressure of the suction device so the gauge on the calibrated negative pressure test gauge reads 150mmHg and read the corresponding reading on the suction device gauge. The gauge on the suction device should be 150mmHg $\pm$ 19mmHg or between 131mmHg – 169mmHg.

If your gauge is out of calibration, you may need a new gauge or a new regulator and gauge assembly (**SSCOR** Part #22012-KIT2314).



4. Check the airflow
 - A. Connect tubing from the patient port of the canister to a calibrated airflow meter. Ensure the regulator gauge is adjusted to the high setting (fully turned clockwise) and turn the device on. The reading on the calibrated airflow meter should be above 30 LPM.
 - B. If the device does not produce 30 LPM check to ensure all tubing connections are tight, the canister lid is securely

fastened to the canister and all the port covers on the canister lid are secure.

- C. If your device still does not produce at least 30 LPM airflow check the airflow at the barb fitting at the top of the device #9. The airflow should be at least 35 LPM at the barb fitting at the top of the device when the device is plugged into AC power.



- D. If the device still does not meet the airflow specification, call SSCOR technical support (1-818-504-4054) to trouble shoot.

After these tests are complete, plug the device in to charge before releasing the device back to the floor. The battery condition LED on the control panel should light green when the device is plugged in and the battery is fully charged.

If your device fails to meet any of these specifications or if you have any questions, call SSCOR technical support at 1-818-504-4054, or you can email techsupportgrp@sscor.com.