

CARDIAC SCIENCE AEDs

G3 *third generation*

Operation and Service Manual

POWER²HEART[®] **AED** **G3**
automated external defibrillator | 9300E

POWER²HEART[®] **AED** **G3**
automated external defibrillator | 9300A

AUT[♥]MATIC



CARDIAC SCIENCE

Limited Warranty

Cardiac Science Corp. ("Cardiac Science") warrants to the original purchaser that its AEDs and stated battery operating life will be free of any defect in material and workmanship according to the terms and conditions of this Limited Warranty ("Limited Warranty"). For purposes of this Limited Warranty, the original purchaser is deemed to be the original end user of the product purchased. This Limited Warranty is NONTRANSFERABLE and UNASSIGNABLE.

For How Long?

Seven (7) years from the date of the original shipment to the original purchaser for Powerheart AED G3 automated external defibrillators and G3 AED battery P/N (9146). Disposable defibrillation pads shall be warranted until the expiration date. Lithium batteries P/N (9146) have a full operational replacement warranty of four (4) years from the date of installation into a Powerheart AED G3. One (1) year from the date of original shipment to the original purchaser for Cardiac Science AED accessories. The terms of the Limited Warranty in effect as of the date of original purchase will apply to any warranty claims.

What You Must Do

Please complete and submit the Warranty Validation Form within 30 days of original shipment located at http://www.cardiacscience.com/products/aed_warranty.cfm. If the purchaser does not have internet access, call (888) 466-8686.

To obtain warranty service for your product, call us toll free at (888) 466-8686 seven days a week, 24 hours a day. Our customer service representative will try to resolve your issue over the phone. If necessary, and at our sole discretion, we will arrange for service or a replacement of our product.

What We Will Do

If your Cardiac Science product is returned within 30 days of the date it was purchased, at the direction of a customer service representative, we will replace it with a new product of equal value at no charge to you, provided the warranty applies.

If your Cardiac Science product is returned, at the direction of a customer service representative, after 30 days but within the warranty period, Cardiac Science, at its sole discretion, will repair your product or replace it. The repaired or replacement product will be warranted subject to the terms and conditions of this Limited Warranty for either (a) 90 days or (b) the remainder of the original warranty period, whichever is longer, provided the warranty applies and the warranty period has not expired.

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This Limited Warranty does not cover defects or damages of any sort resulting from, but not limited to, accidents, damage while in transit to our service location, alterations, unauthorized service, unauthorized product case opening, failure to follow instructions, improper use, abuse, neglect, fire, flood, war or acts of God. Cardiac Science makes no warranty claim as to the compatibility with Cardiac Science products with non Cardiac Science products.

This Limited Warranty is Void if...

- Any Cardiac Science product is serviced or repaired by any person or entity other than Cardiac Science unless specifically authorized by Cardiac Science.
- Any Cardiac Science product case is opened by unauthorized personnel or if a product is used for an unauthorized purpose.
- Any Cardiac Science product is used in conjunction with incompatible parts or accessories, including but not limited to batteries. Parts and accessories are not compatible if they are not Cardiac Science products or the functional equivalent.

If The Warranty Period has Expired...

If your Cardiac Science product is not covered by our Limited Warranty, call us toll free at (888) 466-8686 for advice as to whether we can repair your Cardiac Science product, and for other repair information, including charges. Charges for non-warranty repairs will be assessed and are your responsibility. Upon completion of the repair, the terms and conditions of this Limited Warranty shall apply to such repair or replacement product for a period of 90 days.

This warranty gives you specific legal rights, and you may also have other rights, which vary from state to state.

CAUTION

Federal law restricts this device to be sold by or on the order of a physician or practitioner licensed by state law in which he/she practices to use or order the use of the device.

IMPORTANT

Read this Operation and Service Manual carefully. It contains information about your safety and the safety of others. Become familiar with the controls and how to use the AED properly before operating the product.

Cardiac Science AEDs are manufactured by:

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Bothell, WA 98021 U.S.A.

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PATENTS

This device is covered by the following U.S. and foreign patents:

5,792,190, 5,999,493, 5,402,884, 5,579,919, 5,749,902, 5,645,571, 6,029,085, 5,984,102, 5,919,212, 5,891,172, 5,674,266, 5,700,281, 5,891,173, 5,968,080, 6,263,239, 5,797,969, D402,758, D405,754, 5,909,138, 6,173,203, 6,088,616, 5,897,576, 5,955,956, 6,083,246, 6,064,909, 6,038,473, 5,868,794, 6,115,638, 6,366,809, 5,474,574, 6,246,907, 6,289,243, 6,411,846, 6,480,734, 6,658,290, EP00756878

Other U.S. and foreign patents pending.

LIMITED WARRANTY

The Cardiac Science AED Operation and Service Manual and any and all information contained herein do not constitute any warranty as to the Powerheart AED G3, Powerheart AED G3 Automatic or any related products in any manner whatsoever. The “Limited Warranty” is shipped with the AED and serves as the sole and exclusive warranty provided by Cardiac Science regarding Cardiac Science AED products.

ORDER ENTRY

To order additional Cardiac Science AEDs or accessories:

Worldwide

Toll Free: 800.991.5465

Telephone: 425.402.2690

Fax: 425.402.2001

Email: customerservice@cardiacscience.com

TECHNICAL SUPPORT

To receive 24-hour customer support:

US/INTERNATIONAL

Toll Free: +1.888.466.8686

Or: +1.425.402.2691

Email: techsupport@cardiacscience.com

www.cardiacscience.com

There is no charge to the customer for a customer support call. Please have the serial and model numbers available when contacting Customer Service. (The serial and model numbers are located on the underside of the Cardiac Science AED.)

NOTICE OF RIGHTS

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DEFIBRILLATOR TRACKING

Defibrillator manufacturers and distributors are required, under the Safe Medical Devices Act of 1990, to track the location of defibrillators they sell. Please notify Cardiac Science Customer Service in the event that your defibrillator is sold, donated, lost, stolen, exported, destroyed or if it was not purchased directly from Cardiac Science Corp.

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SECTION 1 – SAFETY

OVERVIEW

This section presents safety information to guard against injury to persons and damage to the Cardiac Science AED G3.

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SAFETY TERMS AND DEFINITIONS

BEFORE OPERATING THE CARDIAC SCIENCE AED G3

Become familiar with the various safety alerts in this section.

Safety alerts identify potential hazards using symbols and words to explain what could potentially harm you, the patient, or the Cardiac Science AED G3.

SAFETY TERMS AND DEFINITIONS

The triangle attention symbol shown below, left, identifies the potential hazard categories. The definition of each category is as follows:



DANGER: This alert identifies hazards that will cause serious personal injury or death.



WARNING: This alert identifies hazards that may cause serious personal injury or death.



CAUTION: This alert identifies hazards that may cause minor personal injury, product damage, or property damage.

PRODUCT MODELS

This manual is for Cardiac Science AED models 9300E and 9300A. They share a basic set of features and differences are noted throughout the manual.

PRODUCT REFERENCES

For purposes of retaining simple, clear instructions in this manual, note the product references used. Features, specifications, operating instructions and maintenance common to product models will be referred to as:

“Powerheart AED”, “AED” or “device”

Refers to Cardiac Science AED G3 models 9300E and 9300A

SAFETY ALERT DESCRIPTIONS

The following is a list of Cardiac Science AED safety alerts that appear in this section and throughout this manual. You must read, understand, and heed these safety alerts before attempting to operate the AED.



DANGER: Fire and Explosion Hazard

Exercise caution when operating the AED close to flammable gases (including concentrated oxygen) to avoid possible explosion or fire hazard.



WARNING: Shock Hazard

Defibrillation shock current flowing through unwanted pathways is potentially a serious electrical shock hazard. To avoid this hazard during defibrillation abide by all of the following:

- Do not touch the patient, unless performance of CPR is indicated
- Do not touch metal objects in contact with the patient
- Keep defibrillation pads clear of other pads or metal parts in contact with patient
- Disconnect all non-defibrillator proof equipment from the patient before defibrillation



WARNING: Shock and Possible Equipment Damage

Disconnect all non-defibrillator proof equipment from the patient before defibrillation to prevent electrical shock and potential damage to the equipment.



WARNING: Electric Shock and Fire Hazard

Do not connect any telephones or unauthorized connectors to the socket on this equipment.



WARNING: Battery is Not Rechargeable

Do not attempt to recharge the battery. Any attempt to recharge the battery may result in an explosion or fire hazard.



WARNING: Shock Hazard

Do not disassemble the AED! Failure to observe this warning can result in personal injury or death. Refer maintenance issues to Cardiac Science authorized service personnel.



CAUTION: Temperature Extremes

Exposing the AED to extreme environmental conditions outside of its operating parameters may compromise the ability of the AED to function properly. The RescueReady® daily self-test verifies the impact of extreme environmental conditions on the AED. If the daily self-test determines environmental conditions outside of the AED's operating parameters, a "Service Required" alert will be issued to prompt the user to move the AED to environmental conditions within the acceptable operating parameters at once. See Section 7 – Technical Data, Parameters, Operation and Standby Conditions.



CAUTION: Lithium Sulfur Dioxide Battery

Pressurized contents: never recharge, short circuit, puncture, deform, or expose to temperatures above 65°C (149°F). Remove the battery when discharged.



CAUTION: Battery Disposal

Recycle or dispose of the lithium battery in accordance with all federal, state and local laws. To avoid fire and explosion hazard, do not burn or incinerate the battery.



CAUTION: Use only Cardiac Science Approved Equipment

Using batteries, pads, cables, or optional equipment other than those approved by Cardiac Science may cause the AED to function improperly during a rescue.



CAUTION: Possible Improper AED Performance

Using pads that are damaged or expired may result in improper AED performance.



CAUTION: Serial Communication Cable

The AED will not function during a rescue when the serial communication cable is connected to its serial port. When the serial communication cable is connected to the AED during a rescue, the prompt "Remove Cable to Continue Rescue" will be heard until you remove the serial communication cable.



CAUTION: Possible Radio Frequency (RF) Susceptibility

RF susceptibility from cellular phones, CB radios and FM 2-way radio may cause incorrect rhythm recognition and subsequent shock advisory. When attempting a rescue using the AED, do not operate wireless radiotelephones within 1 meter of the AED – turn power OFF to the radiotelephone and other like equipment near the incident.



CAUTION: Possible Interference with Implanted Pacemaker

Therapy should not be delayed for patients with implanted pacemakers and a defibrillation attempt should be made if the patient is unconscious and not breathing. The AED has pacemaker detection and rejection, however with some pacemakers the AED may not advise a defibrillation shock.¹

Placing Pads:

- Do not place the pads directly over an implanted device.
- Place the pad at least one inch from any implanted device.



CAUTION: Moving the Patient During a Rescue

During a rescue attempt, excessive jostling or moving of the patient may cause AEDs to improperly analyze the patient's cardiac rhythm. Stop all motion or vibration before attempting a rescue.



CAUTION: Serial Communication Cable

The serial communication cable is only for use with the AED; it is not to be used with a telephone.



CAUTION: Systems Statement

Equipment connected to the analog and digital interfaces must be certified to the respective IEC standards (i.e. IEC 950 for data processing equipment and IEC 601-1 for medical equipment). Furthermore, all configurations shall comply with the system standard IEC 601-1-1. Anybody who connects additional equipment to the signal input part or signal output part configures a medical system, and is therefore, responsible that the system complies with the requirements of the system standard IEC 601-1-1.



CAUTION: Case Cleaning Solutions

When disinfecting the case, use a non-oxidizing disinfectant, such as ammonium salts or a glutaraldehyde based cleaning solution, to avoid damage to the metal connectors.

¹ Cummins, R., ed., *Advanced Cardiac Life Support*; AHA (1994): Ch. 4.

SYMBOL DESCRIPTIONS

The following symbols may appear in this manual, on the AED, or on its optional components. Some of the symbols represent standards and compliances associated with the AED and its use.



Dangerous Voltage: The defibrillator output has high voltage and can present a shock hazard. Please read and understand all safety alerts in this manual before attempting to operate the AED.



Attention!: Identifies important information in this manual, on the AED, or on its component parts regarding the safe and proper use of the AED.



Defibrillator Proof Type BF Equipment: The AED, when connected to the patient's chest by the pads, can withstand the effects of an externally applied defibrillation shock.



CE Mark: This equipment conforms to essential requirements of the Medical Device Directive 93/42/EEC.

IP24 The AED is protected against the effects of splashing water in accordance with IEC 60529.



Classified by ETL Semko with respect to electric shock, fire and mechanical hazards only in accordance with UL 60601-1, CAN/CSA C22.2 No.601.1-M90, EN60601-1 and EN60601-2-4. Conforms to UL Standard UL60601-1. Certified to CAN/CSA Standard C22.2 No. 601.1-M90.



International symbol for ON. Open the lid to turn on the AED.



Open the lid to turn ON the AED.



Indicates the AED battery status. The illuminated areas indicate the remaining battery capacity.



Check pads. The pads are missing, not connected or have compromised functionality.



Indicates AED requires maintenance by authorized service personnel.



When the **SHOCK** indicator is lit, push this button to deliver a defibrillation shock.



When the **CONTINUE** indicator is lit, push this button to clear the internal memory to allow storage of new rescue data in the AED. (Only for models not equipped with Multiple Rescue software)

SYMBOL DESCRIPTIONS (CONT.)



A red indicator with a BLACK X means the AED requires operator attention or maintenance, and is not RescueReady.



A green indicator without a BLACK X means the AED is RescueReady.



Use pads by this date.



Date of manufacture, year and month.



Date of factory recertification (R).



Latex free.



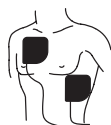
Disposable. Single patient use only.



Tear here to open.



Do not recharge battery.



Position of pads on the chest of patient.



For use by or on the order of a Physician, or persons licensed by state law.



Dispose of properly in accordance with all state, province, and country regulations.



Do not incinerate or expose to open flame.



Explosion hazard: Do not use in the presence of a flammable gas, including concentrated oxygen.



Upper and lower temperature limits.





Device model number, battery model number.





Lot number

 Option number

 Lithium sulfur dioxide

 Serial communication port


 Additional information is provided in the AED Operation and Service Manual.

 Points to important information regarding the use of the AED.

 Lift here

 Manufacturer

 Authorized European Representative

 Symbol for the marking of electrical and electronic equipment that must be recycled.

SECTION 2 – INTRODUCTION

OVERVIEW

This section presents information about the AED, its use, and the training requirements for operation.

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AED DESCRIPTION

The AED is a self-testing, battery-operated automated external defibrillator (AED). After applying the AED's electrodes (pads) to the patient's chest, the AED automatically analyzes the patient's electrocardiogram (ECG) and advises the operator to push the button and deliver a shock if needed. For the Powerheart AED G3 Automatic, the AED automatically delivers a shock if needed. The AED guides the operator through the rescue using a combination of voice prompts, audible alerts, and visible indicators.

INDICATIONS FOR USE

The AED with STAR Biphasic Waveform is intended to be used by personnel who have been trained in its operation. The operator should be qualified by training in basic life support, CPR/AED or other physician-authorized emergency medical response. The device is indicated for emergency treatment of victims exhibiting symptoms of sudden cardiac arrest who are unresponsive and not breathing. If the victim is breathing post-resuscitation, the AED should be left attached to allow for acquisition and detection of the ECG rhythm. If a shockable ventricular tachyarrhythmia recurs, the device will charge automatically and advise the operator to deliver therapy, or for the Powerheart AED G3 Automatic, automatically deliver a shock if needed.

When the patient is a child under 8 years of age or weighs less than 55 lbs (25kg), the AED should be used with the Model 9730 Pediatric Attenuated Defibrillation Pads. Therapy should not be delayed to determine the patient's exact age or weight.

RHYTHMx® AED ECG ANALYSIS ALGORITHM

The RHYTHMx AED ECG analysis algorithm provides ECG detection capabilities. The features available with the AED include the following:

- Detection Rate
- Asystole Threshold
- Noise Detection
- Non-Committed Shock
- Synchronized Shock
- Pacemaker Pulse Rejection
- SVT Discriminators
- Supraventricular Tachycardia (SVT) Rate

DETECTION RATE

All ventricular fibrillation (VF) and ventricular tachycardia (VT) rhythms at or above this rate will be classified as shockable. All rhythms below this rate will be classified as non-shockable. This rate is programmable between 120 bpm (beats per minute) and 240 bpm via MDLink Software by the Medical Director. The default Detection Rate is 160 bpm.

ASYSTOLE THRESHOLD

The asystole peak-to-peak threshold is set at 0.08 mV. ECG rhythms at or below 0.08 mV will be classified as asystole and will not be shockable.

NOISE DETECTION

The AED will detect noise artifact in the ECG. Noise could be introduced by excessive moving of the patient or electronic noise from external sources like cellular and radiotelephones. When noise is detected, the AED will issue the prompt *"Analysis Interrupted. Stop Patient Motion"* to warn the operator. The AED will then proceed to reanalyze the rhythm and continue with the rescue.

NON-COMMITTED SHOCK

After the AED advises a shock, it continues to monitor the patient ECG rhythm. If the patient's rhythm changes to a non-shockable rhythm before the actual shock is delivered, the AED will advise that the rhythm has changed and issue the prompt *"Rhythm Changed. Shock Cancelled."* The AED will override the charge and initiate CPR.

SYNCHRONIZED SHOCK

The AED is designed to automatically attempt to synchronize shock delivery on the R-wave if one is present. If delivery cannot be synchronized within one second, a non-synchronized shock will be delivered.

PACEMAKER PULSE DETECTION

The AED contains pacemaker pulse detection circuitry to detect pulses from an implanted pacemaker.

SVT DISCRIMINATORS

The AED is supplied with the SVT Discriminator enabled and with the default setting “NO THERAPY FOR SVT”. With the factory default setting of “NO THERAPY FOR SVT”, the AED will not shock an SVT rhythm.

SVT Discriminators are sophisticated filters that analyze the morphology of the ECG waveforms and distinguish VF/VT from SVT and Normal Sinus Rhythms (NSR). The SVT Discriminator will only be applied to rhythms that fall between the Detection Rate and the SVT Rate. The factory default setting for this feature is “NO THERAPY FOR SVT”, however the Medical Director can enable this feature using MDLink.

SVT RATE

All rhythms with rates between the Detection Rate and SVT Rate will be screened through a number of SVT Discriminators to classify them into VF/VT or SVT. Rhythms classified as SVT between the two set rates are not shockable. All SVT rhythms above the rates will be classified as shockable. The SVT Rate must be greater than the Detection Rate and is selectable between 160 and 300 bpm or, “NO THERAPY FOR SVT” can be selected via MDLink Software by the Medical Director.

RESCUE PROTOCOL

The AED rescue protocol is consistent with the guidelines recommended by the American Heart Association (AHA)¹ and the International Liaison Committee on Resuscitation (ILCOR).

Upon detecting a shockable cardiac rhythm, the AED advises the operator to press the **SHOCK** button to deliver a shock and then advises the operator to start CPR.

For the Powerheart AED G3 Automatic, upon detecting a shockable rhythm, the AED will automatically deliver a shock and then advise the operator to start CPR.



Note: The standard CPR protocol of 120 seconds can be modified from 60 to 180 seconds in MDLink.

STAR BIPHASIC WAVEFORM



The STAR Biphasic Waveform is designed to measure the patient's impedance and deliver a customized shock. This allows the delivery of an optimized energy level to each patient. The energy levels for the Powerheart AED are available in three different defibrillation shock² configurations. See table on next page and Section 7 for additional information.

STAR BIPHASIC ENERGY PROTOCOLS FOR POWERHEART AED

Cardiac Science's patented STAR[®] Biphasic defibrillation waveform will deliver variable escalating energy that is customized to each patient's needs based upon a patient's thoracic impedance. This customization adjusts for the unique physical differences between patients. The range of impedance over which the device will deliver a shock is 25-180 Ohms. The Powerheart AED comes equipped with five different FDA cleared biphasic energy protocols.

The operator, with guidance, direction and implementation from his/her designated AED program Medical Director, may select from one of these five protocols when placing the Powerheart AED into service. The Powerheart AEDs factory default energy protocol is 200-300-300 Joule (J) escalating Variable Energy (VE). The first shock is delivered within the range of 140J-250J (200J nominal). Subsequent shocks are delivered within a range of 190J-360J (300J nominal). See next page.

¹ "Guidelines 2005 for Cardiopulmonary Resuscitation and Emergency Cardiovascular Care" American Heart Association; *Circulation* Vol 112, Issue 24 Suppl. Dec 13, 2005

² The ultra-low current, low current and high current shocks are variable energy. The actual energy is determined by the patient's impedance.

These protocols are selected by using our MDLink software program. The five biphasic energy protocols available are as follows:

Energy Protocols	Shock Sequence¹	Energy Level	Energy Range (J)
Factory Default	1.	200VE	140J-250J
	2.	300VE	190J-360J
	3.	300VE	190J-360J
Protocol #2	1.	200VE	140J-250J
	2.	200VE	140J-250J
	3.	300VE	190J-360J
Protocol #3	1.	150VE	105J-195J
	2.	200VE	140J-250J
	3.	200VE	140J-250J
Protocol #4	1.	150VE	105J-195J
	2.	150VE	105J-195J
	3.	200VE	140J-250J
Protocol #5	1.	200VE	140J-250J
	2.	200VE	140J-250J
	3.	200VE	140J-250J

¹ The ultra-low current, low current and high current shocks are variable energy. The actual energy is determined by the patient's impedance.

OPERATOR TRAINING REQUIREMENTS

Persons authorized to operate the AED must have all of the following minimum training.

- Defibrillation training and other training as required by state, province, or country regulations.
- Training on operation and use of the AED.
- Additional training as required by the physician or Medical Director.
- A thorough understanding of the procedures in this manual.



Notes: Keep valid certificates of training and certification as required by state, province, or country regulations.

SECTION 3 – GETTING STARTED

OVERVIEW

This section presents information on unpacking and setting up the AED

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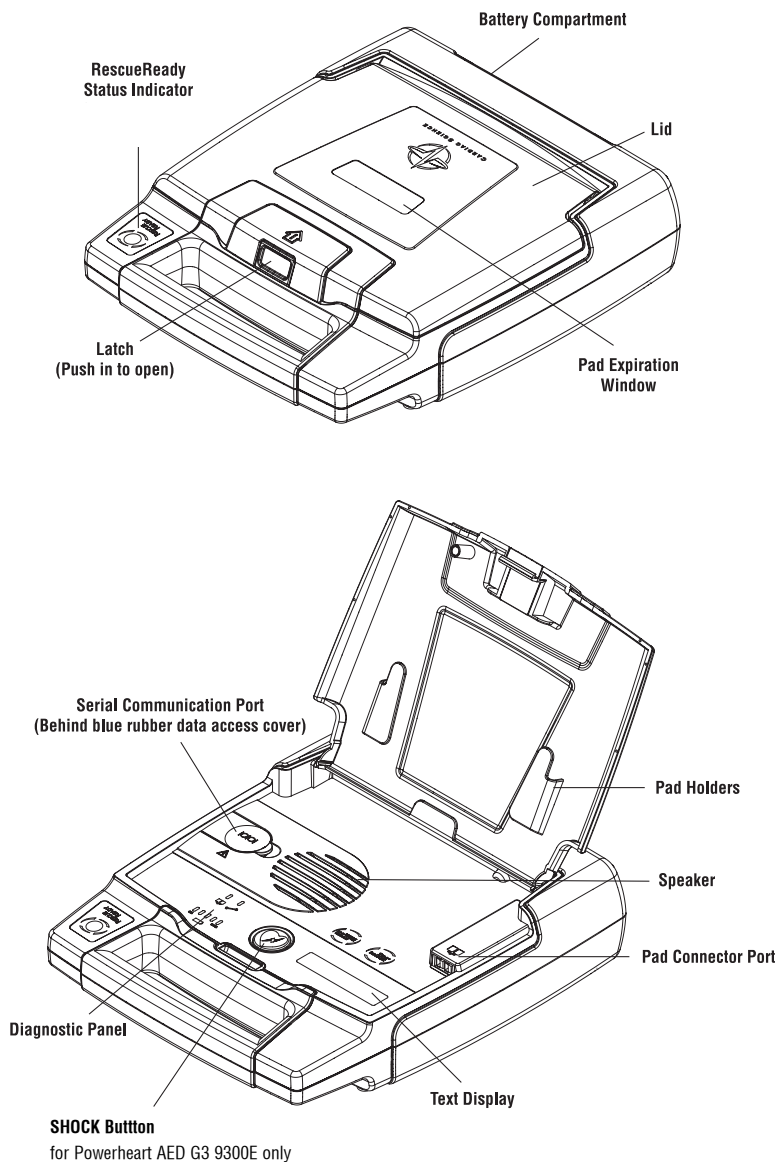
UNPACKING AND INSPECTING

Every attempt is made to ensure your order is accurate and complete. However, to be sure that your order is correct, verify the contents of the box against your packing slip.

If you have any questions about your order, contact Customer Service. See page 4.

AED PARTS

The following drawings show the AED parts and their locations.



AED MODES

Operating Mode: Defined as having the battery installed and the lid open. This is the mode the AED would be in during an actual rescue situation.

Standby Mode: When the battery is installed, but the lid is closed. In this mode the AED is not being used in a rescue. The device will conduct its routine self-tests to ensure proper operation.

Storage Mode: When the battery is removed, such as during shipping or transport. With the battery removed, the AED is unable to perform self-tests or rescues.

ENVIRONMENTAL OPERATING AND STANDBY CONDITIONS

See Section 7 – Technical Data, Parameters, Environmental Operation and Standby Conditions.



CAUTION: Temperature Extremes

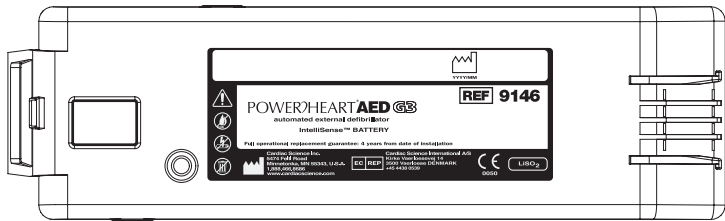
Exposing the AED to extreme environmental conditions outside of its operating parameters may compromise the ability of the AED to function properly. The RescueReady® daily self-test verifies the impact of extreme environmental conditions on the AED. If the daily self-test determines environmental conditions outside of the AED's operating parameters, a "Service Required" alert will be issued to prompt the user to move the AED to environmental conditions within the acceptable operating parameters at once.

SHIPPING AND TRANSPORT CONDITIONS

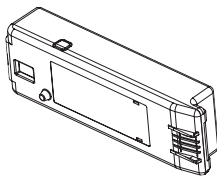
(For up to 1 week)

See Section 7 – Technical Data, Safety and Performance Standards/Shipping and Transportation Conditions.

INTELLISENSE BATTERY



The Cardiac Science IntelliSense battery technology offers you the most advanced battery capabilities available for defibrillators. Cardiac Science IntelliSense batteries contain an integrated memory chip that automatically stores important usage information, enabling the battery to maintain a complete history of its operating life. The actual battery history can be reviewed using the RescueLink software. This history includes:



- Battery Identification
- Battery Type
- Original Date of Installation in an AED
- Number of Charges completed
- Time in Operation (hours:minutes)
- Days of Standby Operation
- Battery Capacity Remaining

BATTERY OPERATING LIFE

The battery operating life depends on the type of battery, actual usage and environmental factors.

The following table represents the expected life of the Powerheart AED when used in Standby Mode.

Model	Estimated Shelf Life (from date of manufacture)	Full Operational Replacement Guarantee (from date of installation)	Typical Shocks
9146 Lithium	5 Years	4 Years	up to 290 shocks

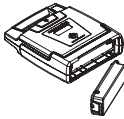
BATTERY SHELF LIFE

The Cardiac Science AED batteries have a shelf life of five years. Shelf life is defined as the length of time a battery can be stored, prior to installation into AED, without degrading its performance.



Note: Storing the battery outside its specific range (0-50°C) will decrease battery life.

BATTERY INSTALLATION



1. With the label on the battery facing the AED battery compartment, insert the battery as shown in the drawing.



2. Push the latched end of the battery firmly into the AED, as shown in the drawing, until the battery snaps into place. The exposed side of the battery should be flush with the outside of the AED case.



3. Open the lid for 5 seconds to initiate a self-test. If the battery is installed properly, the **SMARTGAUGE** battery indicator LEDs will illuminate and the **STATUS INDICATOR** will turn **GREEN**. If service is required, then the **SERVICE** indicator will illuminate; call for service.



WARNING: Battery is Not Rechargeable

Do not attempt to recharge the battery. Any attempt to recharge the battery may result in an explosion or fire hazard.



CAUTION: Lithium Sulfur Dioxide Battery

Pressurized contents: never recharge, short circuit, puncture, deform, or expose to temperatures above 65°C (149°F). Remove the battery when discharged.



CAUTION: Battery Disposal

Recycle or dispose of the lithium battery in accordance with all federal, state and local laws. To avoid fire and explosion hazard, do not burn or incinerate the battery.



CAUTION: Use only Cardiac Science Approved Equipment

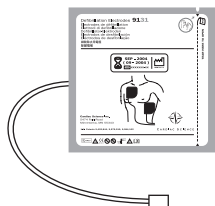
Using batteries, pads, cables, or optional equipment other than those approved by Cardiac Science may cause the AED to function improperly during a rescue.



CAUTION: Possible Improper AED Performance

Using pads that are damaged or expired may result in improper AED performance.

PADS



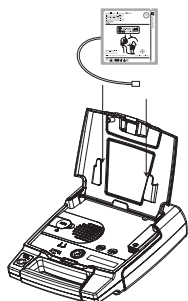
The defibrillation pads come in a ready-to-use, sealed package containing one pair of self-adhesive pads with an attached cable and connector. The pads are disposable and should be discarded after each rescue.

The pads have a limited shelf life and should not be used beyond the expiration date. Keep a fresh, unopened pair of pads plugged into the AED at all times. Refer to the pad package label for operation temperatures.

An audible and visual alert will indicate after the self-test if the pads are missing, unplugged or damaged.

PAD INSTALLATION

1. Open the lid of the AED.
2. Place the pad package into the lid so that the expiration label is visible through the clear window on the lid. The expiration date of the pads will then be readable without opening the lid of the AED.
3. Match the color of the connectors (red to red), then plug the pad connector into the AED case as shown in the drawing. Once the pad connector is plugged into AED, the PAD indicator should extinguish.
4. Tuck the excess cable length in the bottom holder as shown in the drawing. With the pad package completely secured to the AED lid, close the lid.
5. Make sure the expiration date is visible through the clear window of the lid and check to make sure that the **STATUS INDICATOR** is **GREEN**. If the pads are not installed properly, the **STATUS INDICATOR** will be **RED**; call Customer Service for assistance.



CAUTION: Use only Cardiac Science Approved Equipment

Using batteries, pads, cables, or optional equipment other than those approved by Cardiac Science may cause the AED to function improperly during a rescue.



CAUTION: Possible Improper AED Performance

Using pads that are damaged or expired may result in improper AED performance.

DIRECTIONS FOR USE:

1. Do NOT open until ready to use, short term use only.
2. Ensure the skin site is clean and dry.
3. Separate one pad from liner.
4. Place one pad on skin in either location.
5. Peel and place remaining pad.

AED INDICATORS

The following indicators are located on the AED.

RESCUEREADY STATUS INDICATOR



The **STATUS INDICATOR** is located on the Powerheart AED handle.

When this indicator is **GREEN**, the AED is RescueReady. This means the AED self-tests have verified the following:

- Battery has an adequate charge.
- Pads are properly connected to the AED.
- Integrity of the internal circuitry is good.



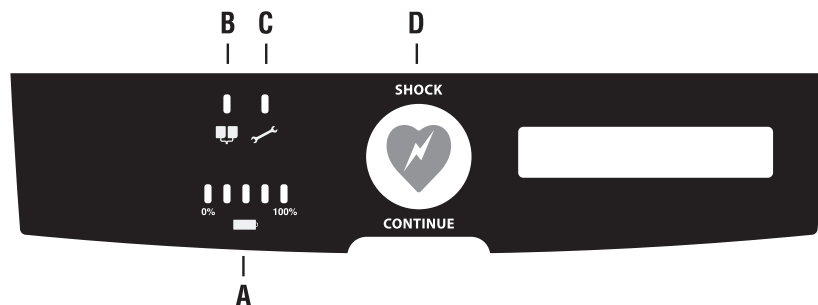
When the **STATUS INDICATOR** is **RED**, maintenance is required.

AUDIBLE MAINTENANCE INDICATOR

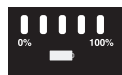
When the daily, weekly or monthly self-test determines service is required, an audible beep is sounded every 30 seconds until the lid is opened or the battery power is depleted. Opening and closing the lid may deactivate the beep. If the error is not corrected by the next automatic self-test, the beep will be reactivated.

DIAGNOSTIC PANEL

- A **SMARTGAUGE BATTERY** Indicator
- B **PADS** Indicator
- C **SERVICE** Indicator
- D **SHOCK** Button (9300E only)



SMARTGAUGE BATTERY STATUS INDICATOR



The **SMARTGAUGE** Battery Status Indicator has five (5) LEDs, four (4) green and one (1) red. The right four green LEDs display the remaining capacity of the battery much like a fuel gauge. With use, the green LEDs gradually go out, from right to left, as battery capacity decreases. When the green LEDs go out and the red LED lights up, replace the battery.



Note: When the red LED initially lights up – upon lid opening or at any time during a rescue – a “*Battery Low*” prompt will be issued at once. However, the AED is capable of delivering at least 9 more defibrillation shocks after the first “*Battery Low*” prompt is issued.

When the AED battery cannot deliver any more shocks, the AED display will show “BATTERY LOW”, the STATUS INDICATOR will be RED, and the device will “beep” every 30 seconds. To continue the rescue, leave the lid open, remove the battery, and replace with a fresh battery. If battery replacement takes longer than 60 seconds, the first rescue will be terminated and a second rescue will begin upon new battery insertion.

PADS INDICATOR



The **PADS** LED lights up when the pads are:

- Not properly connected to the AED.
- Not within operational specifications (cold, dried, damaged).
- Disconnected from the patient during a rescue.

SERVICE INDICATOR



The **SERVICE** LED lights up when the AED requires maintenance that can only be performed by qualified service personnel.

SHOCK INDICATOR



For model 9300E only. The AED has one button called the **SHOCK** button. The word **SHOCK** and the shock button indicator LED will illuminate red when the AED is ready to deliver a defibrillation shock to the patient.

TEXT DISPLAY



The text display has 2 lines of text. The text display provides the operator with information regarding system initialization, text prompts and data during a rescue, and diagnostics.

System initialization occurs when the lid is first opened. The text display shows the operator the identifiers for the internal code, voice prompts and text prompts versions. The text display also shows the current date and time.

During a rescue, the text display shows the number of shocks delivered and the elapsed time from the beginning of the rescue (when the lid was first opened). During CPR, a countdown timer will be displayed. The text version of the voice prompts will also be displayed.



Note: There is a 3 second delay between the time the AED lid is opened and the start of the rescue. This 3 second delay is not included in the elapsed rescue time.

SETTING THE AED INTERNAL CLOCK

The internal clock is preset at Central Standard Time and should be reset to the correct date and local time. If applicable, the AED will automatically adjust itself for Daylight Savings Time. This feature can be turned off using the MDLink software. To set the clock, you will need a Windows 98 or newer PC, RescueLink software installed, and the AED serial cable connected to the PC.

To set the clock settings:

- Ensure that the PC is set at the correct local time and date.
- Open the lid of the AED and run the RescueLink software on the PC.
- Connect the cable to the serial port on the AED.
- Verify that the voice prompt states *"Communications Mode"*.
- Click **COMMUNICATIONS** on the main menu. Select **AED DATE AND TIME**.
- Click on the **GET** button to review the current time in the AED.
- If the time and date is incorrect, click **SET** to set new time and date. The AED date and time will automatically be updated to the PC's time and date.

VOICE PROMPTS AND TEXT DISPLAY

The voice prompts activate when the AED lid is opened and help guide the operator through the rescue. The AED text display provides a visual display of most of the audible voice prompts.

The following table lists the voice and text prompts and a description of when the prompts are issued.

VOICE PROMPT	TEXT DISPLAY	SITUATION
<i>"Tear Open Package and Remove Pads."</i>	TEAR OPEN PACKAGE REMOVE PADS	When the lid is opened, this phrase is repeats twice to initiate the rescue sequence.
<i>"Peel One Pad from Plastic Liner."</i>	PEEL ONE PAD FROM PLASTIC LINER	Repeats until one pad is peeled off of the liner.
<i>"Place One Pad on Bare Upper Chest."</i>	PLACE ONE PAD ON BARE UPPER CHEST	Repeat twice while one pad is placed.
<i>"Peel Second Pad and Place on Bare Lower Chest as Shown."</i>	PEEL SECOND PAD PLACE ON LOWER CHEST	Repeats until both pads are placed on the patient.
<i>"Do Not Touch Patient! Analyzing Rhythm."</i>	DO NOT TOUCH PATIENT ANALYZING RHYTHM	When the AED is analyzing the cardiac rhythm of the patient.
<i>"Shock Advised."</i>	SHOCK ADVISED	When the AED is preparing to deliver a defibrillation shock.
<i>"Charging."</i>	CHARGING	Repeats while AED is charging.

VOICE PROMPT	TEXT DISPLAY	SITUATION
<i>"Stand Clear! Push Flashing Button to Deliver Shock."</i>	<i>STAND CLEAR PUSH BUTTON TO SHOCK</i>	After the AED is fully charged and ready to deliver the defibrillation shock. The RED SHOCK indicator flashes and the phrase repeats for 30 seconds or until the SHOCK button is pushed.
<i>Shock Delivered.</i>	<i>SHOCK DELIVERED</i>	After the AED delivers a defibrillation shock
<i>"It is now safe to touch the patient."</i>	<i>IT IS NOW SAFE TO TOUCH THE PATIENT.</i>	Advises the rescuer when it is safe to touch the patient.
<i>Start CPR</i>	<i>START CPR</i>	After the AED delivers a defibrillation shock After the AED detects a non-shockable rhythm.
<i>Give 30 compressions Then Give Two Breaths</i>	<i>30 COMPRESSIONS 2 BREATHS</i>	Perform CPR for 2 minutes.
<i>"Check Pads"</i>	<i>CHECK PADS</i>	Occurs when patient impedance is too low or too high.
<i>"Battery Low"</i>	<i>BATTERY LOW</i>	Occurs once when the battery voltage becomes low, although a rescue can continue for approximately 9 more shocks. When the battery is too low to do a rescue, the following will occur: 1) "BATTERY LOW" will show on the LCD 2) STATUS INDICATOR will turn RED 3) AED will BEEP once every 30 seconds You must replace the battery before continuing with the rescue. If completely depleted, all AED activity will terminate.
<i>(none)</i>	<i>REMOVE BATTERY COMPLETELY</i>	The AED will show "REMOVE BATTERY COMPLETELY" when the battery is partially removed. But when the battery is at the replace level (showing BATTERY LOW), the REMOVE BATTERY COMPLETELY will not be shown, only the BATTERY LOW.
<i>"Analysis Interrupted. Stop Patient Motion."</i>	<i>ANALYSIS INTERRUPTED STOP PATIENT MOTION</i>	When the AED detects ECG noise artifact, stop moving or touching the patient. Remove other electronic devices within a 5 meter radius.
<i>"Open Lid to Continue Rescue"</i>	<i>OPEN LID TO CONTINUE RESCUE</i>	When the lid is inadvertently closed during a rescue, this prompt will repeat for 15 seconds.
<i>"Rhythm Changed. Shock Cancelled."</i>	<i>RHYTHM CHANGED. SHOCK CANCELLED</i>	When the device is prepared to shock then detects a change in rhythm and therefore cancels the shock.

CARDIAC SCIENCE AEDs



VOICE PROMPT	TEXT DISPLAY	SITUATION
<i>"Continue CPR"</i>	<i>CONTINUE CPR</i>	During CPR mode when enabled, or when a rescue is resumed in CPR mode after being interrupted by the lid closing.
<i>"Remove Cable to Continue Rescue."</i>	<i>REMOVE CABLE</i>	When a serial communication cable is connected to the AED during a rescue, the phrase repeats until the cable is disconnected.
<i>"Communications Mode"</i>	<i>COMMUNICATIONS MODE</i>	When the lid is open and the serial communication cable is plugged into the AED.
<i>(Beep)</i>		One "Beep" occurs in 30-second intervals during CPR when enabled by the MDLink software program, "Beep" occurs when the AED requires maintenance.
<i>"Service Required"</i>	<i>SERVICE REQUIRED</i>	Occurs after the self-tests determine that the AED is not functioning properly. The prompt <i>"Service Required"</i> will be heard when the lid is opened. The red SERVICE indicator will illuminate and <i>"Service Required"</i> will repeat until you close the lid. After closing the lid, an alarm beep will be heard until the battery is removed or becomes completely depleted.

For the Powerheart AED G3 Automatic only:

<i>"Stand Clear! Shock will be Delivered in"</i>	<i>STAND CLEAR. SHOCK IN</i>	After the AED is fully charged and ready to deliver the defibrillation shock. The Shock will automatically be administered three seconds after the end of the voice prompt.
<i>"Three, two, one"</i>	<i>THREE TWO ONE</i>	Counts down seconds until a shock is automatically delivered.
<i>"Shock delivered."</i>	<i>SHOCK DELIVERED</i>	After a shock is automatically delivered.
<i>"It is now safe to touch the patient."</i>	<i>IT IS NOW SAFE TO TOUCH THE PATIENT.</i>	Advises the rescuer when it is safe to touch the patient.

SECTION 4 – INSTRUCTIONS FOR USE

OVERVIEW

This section presents information about how to use the AED to perform a rescue.

Topic	Page #
Step 1: Assessment and Pad Placement	34
Step 2: ECG Analysis	35
Step 3: Shock Delivery and CPR Mode	36
Step 4: Post Rescue	37
Warnings	37

STEP 1: ASSESSMENT AND PAD PLACEMENT

PREPARATION

Determine that the patient is over 8 years of age or weighs more than 55 pounds (25 kg) and exhibits the following:

The patient is unresponsive, and
the patient is not breathing.

Remove clothing from the patient's chest. Ensure the skin site is clean and dry. Dry the patient's chest and shave excessive hair if necessary.

Open the AED lid and wait until the LEDs are lit.



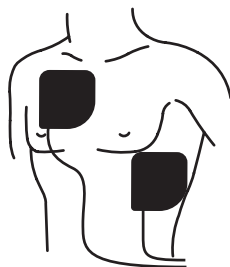
Note: When the patient is a child under 8 years of age or weighs less than 55 lbs (25kg), the AED should be used with the Model 9730 Pediatric Attenuated Defibrillation Pads. Therapy should not be delayed to determine the patient's exact age or weight. See the directions for use accompanying pediatric pads for procedure on changing adult pads to pediatric.

PLACE PADS

The AED will issue the prompt *"Tear Open Package and Remove Pads."* Keep the pads connected to the AED, tear the pad package along the dotted line and remove the pads from the package. Leave the package attached to the pad wires.



After the prompt *"Peel One Pad From Plastic Liner,"* with a firm, steady pull, carefully peel one pad away from the plastic liner.



Then, after the prompt *"Place One Pad on Bare Upper Chest,"* place the pad with the sticky side of on the patient's skin on the upper right chest, placing the top of the pad on the collarbone. Avoid placing the pad directly over the sternum.

Finally, after the prompt *"Peel Second Pad and Place on Bare Lower Chest As Shown,"* pull the second pad from the plastic liner and place it on lower left chest, below and left of the breast.



Note: Cardiac Science's defibrillation pads are non-polarized and can be placed in either position as shown on the pad package.

When the pads are placed, the voice prompt will say *“Do not touch patient. Analyzing Rhythm.”* If the pads are not properly placed or become disconnected at any time during the rescue, the voice prompt *“Check Pads”* will be heard. When this occurs, ensure that:

Pads are firmly placed on clean, dry skin

Pad cable is securely plugged into the AED

STEP 2: ECG ANALYSIS

As soon as the AED detects proper pad placement, the voice prompt *“Do Not Touch Patient. Analyzing Rhythm”* will be heard. The AED will begin to analyze the cardiac rhythm of the patient.

If a shock is advised, the voice prompt will say, *“Shock Advised. Charging.”*

For the Powerheart AED G3 Automatic:

The voice prompt, *“Stand Clear! Shock will be delivered in 3, 2, 1.”* will be heard.

When the AED is charged, it continues to analyze the patient's heart rhythm. If the rhythm changes and a shock is no longer needed, the AED will issue the prompt *“Rhythm Changed. Shock Cancelled,”* disarm and initiate CPR.

If no shock is advised, the AED will prompt to start CPR.

If noise is detected during analysis, the AED will warn you with the prompt *“Analysis Interrupted. Stop Patient Motion”* and restart the analysis. This usually occurs if the patient is excessively jostled or there is a strong electromagnetic emitting electronic device nearby (within 5 meters). Remove the electronic device or stop the excessive motion when you hear this prompt.

STEP 3: SHOCK DELIVERY AND CPR MODE

When the AED is ready to deliver a defibrillation shock, the **SHOCK** button will flash and the prompt *“Stand Clear. Push Flashing Button to Deliver Shock”* will be heard.

Make sure no one is touching the patient and push the **SHOCK** button to deliver a defibrillation shock. If you do not push the **SHOCK** button within 30 seconds of hearing the prompt, the AED will prompt *“It is safe to touch the patient”*. The AED will then prompt you to start CPR.

For the Powerheart AED G3 Automatic ONLY:

When the AED is ready to deliver a shock, the voice prompt, *“Stand Clear! Shock will be delivered in 3, 2, 1.”* will be heard then the AED will deliver a shock.

After the AED delivers a defibrillation shock, the voice prompt will say *“Shock Delivered.”* The AED will then prompt you to start CPR.



Note: During a rescue, the text screen displays voice prompts, elapsed time of rescue and number of shocks delivered.

CPR MODE



After shock delivery or detection of a non-shockable rhythm, the AED automatically enters CPR mode. The voice prompt will say, *“It is now safe to touch the patient. Start CPR.”*

During the CPR time-out period. The AED will not interrupt the CPR mode if the patient’s condition changes and the AED detects a shockable rhythm. After the CPR time-out period has expired, the voice prompt *“Do Not Touch Patient. Analyzing Rhythm.”* will be heard.

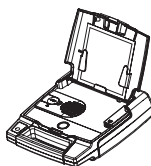


Note: During CPR mode, the text screen displays a countdown timer.

If the patient is conscious and breathing normally, leave the pads on the patient's chest connected to the AED. Make the patient as comfortable as possible and wait for Advanced Life Support [ALS] personnel to arrive. Continue to follow the voice prompts until the ALS personnel arrive, or proceed as recommended by the Medical Director.

STEP 4: POST RESCUE

After transferring the patient to ALS personnel, prepare the AED for the next rescue:



1. Retrieve the rescue data stored in the internal memory of the AED by using RescueLink software installed on a PC (see detailed procedure in the Data Management section).
2. Connect a new pair of pads to the AED.
3. Close the lid.
4. Verify that the **STATUS INDICATOR** on the handle is **GREEN**,

WARNINGS

The following cautions must be observed to prevent problems during the rescue.



DANGER: Fire and Explosion Hazard

Exercise caution when operating the AED close to flammable gases (including concentrated oxygen) to avoid possible explosion or fire hazard.



WARNING: Shock Hazard

Defibrillation shock current flowing through unwanted pathways is potentially a serious electrical shock hazard. To avoid this hazard during defibrillation abide by all of the following:

- Do not touch the patient, unless performance of CPR is indicated
- Do not touch metal objects in contact with the patient
- Keep defibrillation pads clear of other pads or metal parts in contact with patient
- Disconnect all non-defibrillator proof equipment from the patient before defibrillation



WARNING: Shock and Possible Equipment Damage

Disconnect all non-defibrillator proof equipment from the patient before defibrillation to prevent electrical shock and potential damage to the equipment.



WARNING: Electric Shock and Fire Hazard

Do not connect any telephones or unauthorized connectors to the socket on this equipment.



CAUTION: Use only Cardiac Science Approved Equipment

Using batteries, pads, cables, or optional equipment other than those approved by Cardiac Science may cause the AED to function improperly during a rescue.



CAUTION: Possible Improper AED Performance

Using pads that are damaged or expired may result in improper AED performance.



CAUTION: Serial Communication Cable

The AED will not function during a rescue when the serial communication cable is connected to its serial port. When the serial communication cable is connected to the AED during a rescue, the prompt *“Remove cable to continue rescue”* will be heard until you remove the serial communication cable from the AED.



CAUTION: Possible Radio Frequency (RF) Susceptibility

RF susceptibility from cellular phones, CB radios and FM 2-way radio may cause incorrect rhythm recognition and subsequent shock advisory. When attempting a rescue using the AED, do not operate wireless radiotelephones within 1 meter of the AED – turn power OFF to the radiotelephone and other like equipment near the incident.



CAUTION: Possible Interference with Implanted Pacemaker

Therapy should not be delayed for patients with implanted pacemakers and a defibrillation attempt should be made if the patient is unconscious and not breathing. The AED has pacemaker detection and rejection, however with some pacemakers the AED may not advise a defibrillation shock.¹

Placing Pads:

- Do not place the pads directly over an implanted device.
- Place the pad at least an inch from any implanted device.



CAUTION: Moving the Patient During a Rescue

During a rescue attempt, excessive jostling or moving of the patient may cause AEDs to improperly analyze the patient's cardiac rhythm. Stop all motion or vibration before attempting a rescue.

¹ Cummins, R., ed., *Advanced Cardiac Life Support*; AHA (1994): Ch. 4.

SECTION 5 – DATA MANAGEMENT

OVERVIEW

The AED is designed for ease of data management and review. The data stored in internal memory can be displayed on the PC screen using the RescueLink software.

Topic	Page #
Recording Rescue Data	39
Reviewing Rescue Data	39

RECORDING RESCUE DATA

RECORDING DATA IN INTERNAL MEMORY

The AED automatically stores up to 60 minutes of rescue data.

REVIEWING RESCUE DATA

RETRIEVING DATA FROM MEMORY

1. Open the AED lid.
2. Connect the serial cable to the PC and to the AED's serial port under the blue rubber data access cover. The voice prompt will say "*Communications Mode.*"
3. Run the RescueLink software program.
4. Select COMMUNICATIONS, GET RESCUE DATA. On the RescueLink software program.
5. Select INTERNAL MEMORY OF AED then select OK.
6. Select a rescue by clicking on the date and press OK.



WARNING: Electric Shock and Fire Hazard

Do not connect any telephones or unauthorized connectors to the socket on this equipment.



CAUTION: Serial Communication Cable

The serial communication cable is only for use with the AED; it is not to be used with a telephone.

MULTIPLE RESCUE FUNCTIONALITY

The AED can store up to 60 minutes of ECG monitoring time in the AED's internal memory. Multiple rescues can be stored in the internal memory, allowing the rescuer to administer additional rescues without downloading the data to a PC. Should the internal memory become full, the AED will purge rescues as needed, beginning with the oldest rescue.

When downloading data, RescueLink will enable the user to select which rescue to download. See the RescueLink application HELP files for more information.

SECTION 6 – MAINTENANCE & TROUBLESHOOTING

OVERVIEW

This section presents information about the AED diagnostics self-tests, maintenance, and service indications.

Topic	Page #
Self-Tests	41
Indicator Troubleshooting Table	42
Scheduled Maintenance	43
Authorized Repair Service	44
Frequently Asked Questions	45

SELF-TESTS

The AED has a comprehensive self-test system that automatically tests the electronics, battery, pads, and high voltage circuitry. Self-tests are also activated every time you open and close the AED lid.

When performing the self-tests, the AED completes the following steps automatically.





- Turns itself ON, and the **STATUS INDICATOR** changes to **RED**.
- Performs the self-test.
- If successful, the **STATUS INDICATOR** reverts to **GREEN**.
- Turns itself OFF if the lid is closed.

There are three types of automatic self-tests. The Daily Self-test checks the battery, pads, and the electronic components. The Weekly Self-test completes a partial charge of the high voltage electronics current in addition to the items tested in the Daily Self-test. During the Monthly Self-test, the high voltage electronics are charged to full energy.

Self-tests will be initiated upon opening the lid and again upon closing the lid. If the self-test detects an error, the **STATUS INDICATOR** will remain **RED**. Upon closing the lid, an audible alert will be issued. The Diagnostic Panel under the lid will indicate the source of the problem according to the Indicator Troubleshooting Guide Table on the next page.

INDICATOR TROUBLESHOOTING TABLE

The following is a troubleshooting table for the AED indicators.

VIEW	SYMPTOM	SOLUTION
	Red SERVICE indicator (LED) is lit.	Maintenance by authorized service personnel is required. Call Cardiac Science Customer Service (see page 4) or your local Cardiac Science distributor.
	Red PADS indicator (LED) is lit.	Connect the pads or replace with a new pair.
	The last battery indicator (LED) is red.	The battery is low. Replace with a new battery.
	STATUS INDICATOR is RED , and no other indicators on the diagnostic panel are lit.	The battery power is completely depleted. Replace with a new battery. If STATUS INDICATOR remains RED call Cardiac Science Customer Service or your local Cardiac Science distributor.



CAUTION: Temperature Extremes

Exposing the AED to extreme environmental conditions outside of its operating parameters may compromise the ability of the AED to function properly. The RescueReady® daily self-test verifies the impact of extreme environmental conditions on the AED. If the daily self-test determines environmental conditions outside of the AED’s operating parameters, a “SERVICE REQUIRED” alert will be issued to prompt the user to move the AED to environmental conditions within the acceptable operating parameters at once. See Section 7 – Technical Data, Parameters, Operation and Standby Conditions.

SCHEDULED MAINTENANCE

DAILY MAINTENANCE



Check the **STATUS INDICATOR** to ensure that it is **GREEN**. When the indicator is **GREEN**, the AED is ready for a rescue. If the indicator is **RED**, refer to the Troubleshooting Table in this chapter.

MONTHLY MAINTENANCE

1. Open the AED lid.
2. Wait for the AED to indicate status:
Observe the change of the **STATUS INDICATOR** to **RED**.
After approximately 5 seconds, verify that the **STATUS INDICATOR** returns to **GREEN**.
3. Check the expiration date on the electrodes.
4. Listen for the voice prompts.
5. Close the lid and confirm that **STATUS INDICATOR** remains **GREEN**.

ANNUAL MAINTENANCE

Perform the following tests annually to confirm that the diagnostics are functioning properly and to verify the integrity of the case.

Check the Integrity of the Pads and Circuitry



1. Open the AED lid.
2. Remove the pads.
3. Close the lid.
4. Confirm that the **STATUS INDICATOR** turns red.
5. Open the lid and confirm that the **PAD** indicator is lit.
6. Reconnect the pads and close the lid.
7. Make sure the expiration date is visible through the clear window of the lid.
 - a. Check to make sure that the **STATUS INDICATOR** is **GREEN**. If the pads are not installed properly, the **PAD** indicator will illuminate; call Customer Service for assistance.
8. Open the lid and confirm that no diagnostic indicators are lit.
9. Check the expiration date of the pads; if expired, replace them.
10. Check the pads packaging integrity.
11. Close the lid.

Check the Integrity of the Service Indicator (LED) and Circuitry



1. Immediately after opening the AED lid, press and hold the **SHOCK** button and confirm that the **SERVICE** LED is lit.
2. Release the **SHOCK** button.
3. Close the lid.
4. Verify that the **STATUS INDICATOR** remains red.
5. Open the lid and confirm that no diagnostic indicators are lit.
6. Close the lid.
7. Verify that the **STATUS INDICATOR** turns green.

Check the Integrity of the Case

Examine the molded case of the AED for any visible signs of stress. If the case shows signs of stress, contact Cardiac Science Customer Service (See page 4) or contact your local Cardiac Science distributor.



CAUTION: Case Cleaning Solutions

When disinfecting the case, use a non-oxidizing disinfectant, such as ammonium salts or a glutaraldehyde based cleaning solution, to avoid damage to the metal connectors.

AUTHORIZED REPAIR SERVICE

The AED has no user-serviceable internal components. Try to resolve any maintenance issues with the AED by using the Troubleshooting Table presented in this chapter. If you are unable to resolve the problem, contact Cardiac Science Customer Service (See page 4) or contact your local Cardiac Science distributor.



WARNING: Shock Hazard

Do not disassemble the AED! Failure to observe this warning can result in personal injury or death. Refer maintenance issues to Cardiac Science authorized service personnel.



Note: The warranty will be void upon unauthorized disassembly or service of the AED.

FREQUENTLY ASKED QUESTIONS

QUESTIONS AND ANSWERS

1. Q: *Can I give CPR while the AED is analyzing?*
A: No. As with all AEDs, the operator should stop CPR compressions during the analysis phase.
2. Q: *Can I transport the victim while the AED is analyzing?*
A: No. Vehicle motion may cause noise artifacts that could interfere with proper cardiac rhythm analysis. Stop the vehicle when cardiac rhythm analysis is necessary.
3. Q: *Do I need to prepare the chest prior to pad application?*
A: Special preparation is not usually necessary. The chest should be as clean, dry, and as oil free as possible. Follow your Medical Director's instruction.
4. Q: *What happens if the battery is low when I begin a rescue?*
A: When the battery indicator is red, the AED issues a "Battery Low" prompt once; however, the AED is still capable of delivering approximately 9 more defibrillation shocks.

When the AED is not capable of delivering any more shocks, it "beeps" once every 30 seconds. To continue the rescue attempt, leave the lid open and replace the battery. When the battery replacement takes longer than 60 seconds, the first rescue is terminated and the AED will begin to record the events from then on as a separate rescue.

5. Q: *How do I set the AED internal clock?*
A: Set the clock by using the RescueLink Software Program and a PC. See Setting the AED Internal Clock in Chapter 3.
6. Q: *What happens if I close the lid in the middle of a rescue attempt?*
A: If you close the lid during a rescue, you must re-open the lid within 15 seconds to continue the rescue. You will hear the prompt, "Open Lid to Continue Rescue." If the lid remains closed for more than 15 seconds, a new rescue will initiate when the lid is reopened.



Note: If the lid is closed during a rescue while the pads are connected to the patient, the **STATUS INDICATOR** may turn **RED**. When the lid is reopened, however, the rescue may be continued even though the **STATUS INDICATOR** remains **RED**.

7. Q: *My AED is sounding an audible alert. Why? How do I stop it?*
A: The audible alert indicates that the self-test detected a need for maintenance or corrective action. Determine the maintenance required by using the Troubleshooting Table in this chapter. Opening and closing the lid may turn OFF the audible alert until the next self-test. However, the **STATUS INDICATOR** will remain **RED**.
8. Q: *The AED did not sound an audible alert when I removed the pads and closed the lid. Why?*
A: The lid-closed pad self-test only activates the **STATUS INDICATOR**.
The AED allows time for replacement of the pads – as removing pads is a normal procedure after a rescue – or a battery during the post rescue procedure, however, an audible maintenance indicator will be triggered after the next Daily Self-test.
9. Q: *What if I have to perform a rescue in an isolated area and at subzero temperatures?*
A: When travel to a rescue involves exposing the AED to extremely cold temperatures for an extended period of time, keep the pads and the battery warm.

SECTION 7 – TECHNICAL DATA

OVERVIEW

This section presents technical data about the AED.

Topic	Page #
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STAR Biphasic Waveform	52
RHYTHMx ECG Analysis Performance	54

PARAMETERS

OPERATION

- Semi-Automatic (shock advisory)
- Automatic

AUDIBLE ALERTS

- Voice Prompt
- Maintenance Alert

VISIBLE INDICATORS

- Status Indicator
- Battery Status Indicator
- Service Indicator
- Pads Indicator
- Text Display

RESCUE DATA STORAGE

Storage	Capacity
Internal	60 minutes ECG data with event annotation

DIMENSIONS

Measurement	Dimension
Height	8 cm (3.3 in)
Width	27 cm (10.6 in)
Depth	31 cm (12.4 in)

WEIGHT

Model	Weight with Batteries and Pads
9300	3.10 kg (6.6 lb)

ENVIRONMENTAL OPERATION AND STANDBY CONDITIONS

Atmosphere	Condition
Temperature	0°C to 50°C (32°F to 122°F)
Humidity	5% to 95% (non-condensing)
Pressure	57kPa (+15,000ft) to 103kPa (-500ft)

SHIPMENT AND TRANSPORT ENVIRONMENTAL CONDITIONS (for up to 1 week)

Atmosphere	Condition
Temperature	-30°C to 65°C (-22°F to 149°F)
Humidity	5% to 95% (non-condensing)
Pressure	57kPa (+15,000ft) to 103kPa (-500ft)

PADS

- Self-adhesive, disposable defibrillation pads
- Minimum combined surface area: 228cm²
- Extended length of lead wire: 1.3m

LITHIUM BATTERY SPECIFICATIONS

- Output voltage: 12VDC (max)
- Batteries are non-rechargeable
- Lithium contents: 9.2g (max)
- Check local regulations for disposal information

Model	Estimated Shelf Life (from date of manufacture)	Full Operational Replacement Guarantee (from date of installation)	Typical Shocks
9146 Lithium	5 Years	4 Years	up to 290 shocks

The battery operating life depends on the type of battery, actual usage and environmental factors.

BATTERIES AND CAPACITOR CHARGE TIMES

A new battery typically takes 10 seconds to charge the AED to maximum energy.

A battery with reduced capacity causes the red LED light to initially turn ON and typically takes 13 seconds to charge a fully discharged AED to maximum energy.

The maximum time from “Power On” to “Ready to Shock” is 28 seconds for a new rescue.
The maximum time from “Analyze” to “Ready to Shock” is 22 seconds for a new rescue.

AED SELF-TEST SEQUENCE


Frequency of Self-Test	What is Tested?
Daily	Battery, pads, internal electronics, SHOCK button, and software (no charge).
Weekly	Battery, pads, internal electronics, SHOCK button, and software (partial charge).
Monthly (every 28 days)	Battery under load, pads, internal electronics, full-energy charge cycle, SHOCK , and software (full charge).
Open Lid (when lid is opened)	Battery, pads, internal electronics, SHOCK button, and software.
Close Lid (when lid is closed)	Battery, pads, internal electronics, SHOCK button, and software.

SAFETY AND PERFORMANCE STANDARDS

AED MODELS 9300

The AED has been designed and manufactured to conform to the highest standards of safety and performance including electromagnetic compatibility (EMC). The Cardiac Science AED Models 9300 and pads conform to the applicable requirements of the following:

 **CE**
CE Marked by BSI 0086 per the Medical Device Directive 93/42/EEC of European Nations

 **ETL**
Classified by ETL Semko with respect to electric shock, fire and mechanical hazards only in accordance with UL 60601-1, CAN/CSA C22.2 No.601.1-M90, EN60601-1 and EN60601-2-4. Conforms to UL Standard UL60601-1. Certified to CAN/CSA Standard C22.2 No. 601.1-M90.

Electrical, Construction, Safety and Performance

IEC 60601-1 (1998), Amendments 1 (1991) & 2 (1995)
IEC 60601-2-4 (2002)
ANSI/AAMI DF-39 (1993)

Electromagnetic Compatibility (EMC)

IEC 60601-1-2 (2001)
IEC 60601-2-4 Section 36
ANSI/AAMI DF-39 (1993) Section 3.3.21

EMISSIONS

Field	Standard or Compliance
E-M	EN 55011/CISPR 11, Group 1, Class B
Magnetic	ANSI/AAMI DF39, <0.5mT on surface, except for within 5cm of the lid magnet and the speaker

IMMUNITY

Field	Standard or Compliance
E-M	IEC 61000-4-3, Level X, (20V/m) IEC 60601-2-4, Section 36.202.3 (20V/m) AAMI DF39, Section 3.3.21.2.1
Magnetic	IEC 61000-4-8 (2001) IEC 60601-2-4 (2002), Section 36.202.8 AAMI DF39, Section 3.3.21.2.3 80A/m, 47.5Hz – 1,320Hz
ESD	IEC 61000-4-2, Level 3 IEC 60601-2-4 (2002), Section 36.202.2 6KV contact discharge, 8KV air gap discharge

ENVIRONMENTAL CONDITIONS

Condition	Standard or Compliance
Free Fall Drop	IEC 60068-2-32 (1975) AM 2 (1990), 1 meter
Bump	IEC 60068-2-29 (1987), 40g and 6000 bumps
Vibration (Random)	IEC 60068-2-64 (1993): 10Hz – 2KHz, 0.005 – 0.0012 g ² /Hz
Vibration (Sine)	IEC 60068-2-6 (1995): 10Hz – 60Hz, 0.15 mm and 60Hz – 150Hz, 2g
Enclosure Protection	IEC 60529 (2001), IP24

SHIPPING AND TRANSPORTATION CONDITIONS

ISTA Procedure 2A

STAR BIPHASIC WAVEFORM

The waveform generated by the Cardiac Science AED is a BIPHASIC TRUNCATED EXPONENTIAL waveform that is compliant with ANSI/AAMI DF2 and DF39. The following is a graph of the waveform voltage as a function of time when the AED is connected to a 50 Ohm resistive load.

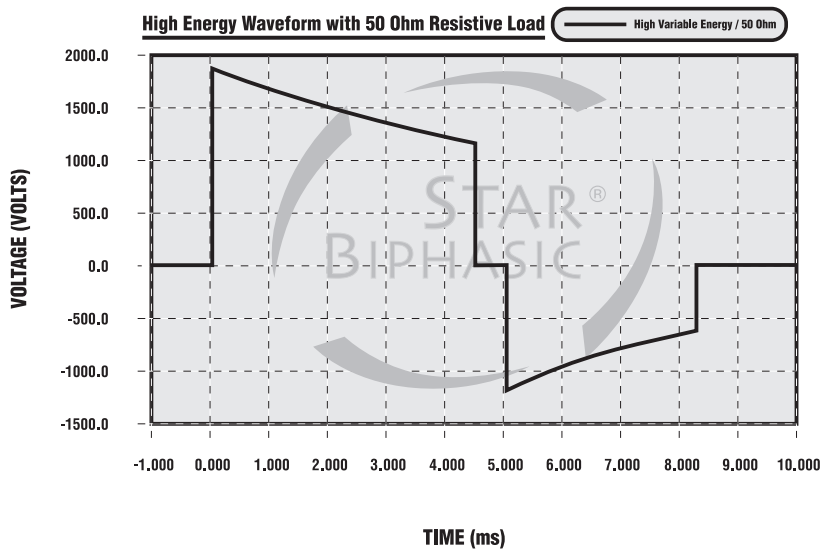


FIGURE A1. STAR BIPHASIC WAVEFORM.

Table A1 - Ultra-low Current Powerheart AED Models 9300 Waveform (all values are typical)

Patient's Impedance (Ohms)	Phase 1		Phase 2		Energy (Joules)
	Voltage (Volts)	Duration (ms)	Voltage (Volts)	Duration (ms)	
25	1390	3.3	730	3.2	145-195
50	1420	4.5	915	3.2	130-175
75	1430	5.8	980	3.2	120-160
100	1435	7.0	1020	3.2	110-150
125	1440	8.3	1040	3.2	105-140

Table A2 - Low Variable Energy Powerheart AED Models 9300 Waveform (all values are typical)

Patient's Impedance (Ohms)	Phase 1		Phase 2		Energy (Joules)
	Voltage (Volts)	Duration (ms)	Voltage (Volts)	Duration (ms)	
25	1570	3.3	825	3.2	200-250
50	1600	4.5	1030	3.2	170-210
75	1610	5.8	1105	3.2	120-160
100	1615	7.0	1150	3.2	150-180
125	1620	8.3	1170	3.2	140-170

Table A3 - High Variable Energy Powerheart AED Models 9300 Waveform (all values are typical)

Patient's Impedance (Ohms)	Phase 1		Phase 2		Energy (Joules)
	Voltage (Volts)	Duration (ms)	Voltage (Volts)	Duration (ms)	
25	1885	3.3	990	3.2	265-360
50	1920	4.5	1240	3.2	235-320
75	1930	5.8	1325	3.2	215-295
100	1940	7.0	1380	3.2	200-270
125	1945	8.3	1405	3.2	190-260

ENERGY LEVELS AND PATIENT IMPEDANCE

The Cardiac Science Biphasic Truncated Exponential (BTE) waveform utilizes variable energy. The actual energy delivered will vary with the patient's impedance and the device will deliver a shock when impedance is between 25-180 Ohms. Energy will be delivered at three different levels referred to as ultra-low variable energy, low variable energy, and high variable energy as shown in the above waveform tables.

RHYTHMx ECG ANALYSIS PERFORMANCE

The AED RHYTHMx ECG Analysis system analyzes the patient’s ECG and advises you when the AED detects a shockable or non-shockable rhythm.

This system makes it possible for a person, with no training in the interpretation of ECG rhythms, to offer defibrillation therapy to victims of sudden cardiac arrest.

CARDIAC RHYTHMS USED TO TEST THE RHYTHM RECOGNITION DETECTION SYSTEM FOR CARDIAC SCIENCE AED

Rhythm Class	Specifications
Shockable Rhythm – VF	Meets AAMI DF 39 requirement and AHA recommendation of Sensitivity ^a of >90%
Shockable Rhythm – VT	Meets AAMI DF 39 requirement and AHA recommendation of Sensitivity of >75%
Non-shockable Rhythm – NSR	Meets AAMI DF 39 requirement (>95%) and AHA recommendation (>99%) of Specificity
Non-shockable – Asystole	Meets AAMI DF 39 requirement and AHA recommendation of Specificity of >95%
Non-shockable – all other rhythms	Meets AAMI DF 39 requirement and AHA recommendation of Specificity of >95%

^a Automatic External Defibrillators for Public Access Defibrillation: Recommendations for Specifying and Reporting Arrhythmia Analysis Algorithm Performance, Incorporating New Waveforms and Enhancing Safety, AHA AED Task Force and approved by the AHA Science Advisory and Coordinating Committee. *Circulation*, 1997(95), pp 1677-1682



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