



# HeartSine samaritan® PAD SAM 360P



User Manual

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## Symbols used in this manual



**Warning: Risk of death or serious injury**



**Caution: Risk of injury**



**Notice: Risk of damage to data or material**



**Further information**

## Symbols used on this device



On/Off

**IP56**

Ingress protection classified as IP56 according to EN 60529



Consult operating instructions



Single use item. Do not re-use



Defibrillation protected,  
Type BF connection



Do not expose to high heat or open flame. Do not incinerate



Does not contain natural rubber latex



Non-sterile



Recyclable



Non-rechargeable battery



Do not short circuit battery



Do not crush battery



Temperature limitation as indicated



Use by yyyy/mm



Dispose of in accordance with country requirements



Automated External Defibrillator

With respect to electrical shock, fire and mechanical hazards only in accordance with

- ANSI/AAMI ES60601-1:2005
- CSA C22.2 NO. 60601-1:2008
- IEC60601-2-4:2010



Follow instructions for use



e.g. "yyE01234567"  
yy = year of manufacture

# Indications for Use

## Indications for use

The HeartSine® samaritan® PAD 360P is indicated for use on victims of cardiac arrest who are exhibiting the following signs:

- Unconscious
- Not breathing
- Without circulation

The samaritan PAD 360P (SAM 360P) is indicated for use on patients greater than 8 years old or over 25 kg (55 lbs) when used with the adult samaritan Pad-Pak™. The SAM 360P is indicated for use on children between 1 and 8 years of age or up to 25 kg (55 lbs) when used with the samaritan Paediatric-Pak™.

## Contraindications for use

If the patient is responsive or conscious, do not use the SAM 360P to provide treatment.

## Intended users

The SAM 360P is intended for use by personnel who have been trained in its operation. Users should have received training in basic life support / AED, advanced life support or a physician-authorised emergency medical response training program.

# Warnings and Cautions



## Warning

### Patients suitable for treatment

The SAM 360P has been designed to work on unconscious, nonresponsive patients. If the patient is responsive or conscious, do not use the SAM 360P to provide treatment.

The SAM 360P uses an interchangeable battery and electrode pack called Pad-Pak. The SAM 360P in combination with an adult Pad-Pak is suitable for use on patients of over 25 kilograms (55 pounds) in weight or equivalent to a child of approximately eight years old or over.

For use on smaller children (from 1 to 8 years old), remove the adult Pad-Pak and install a Paediatric-Pak. If a Paediatric-Pak or an alternative suitable defibrillator is not available, you may use an adult system.

Do not delay treatment trying to find out the patient's exact age and weight.

### Risk of electric shock

The SAM 360P delivers therapeutic electrical shocks that can cause serious harm to either operators or bystanders. Take care to ensure that nobody touches the patient when a shock is to be delivered.

### Avoid opening or repairing

The SAM 360P has no serviceable parts. Do NOT open or repair the device under any circumstances as there could be danger of electric shock. If damage is suspected, replace the SAM 360P immediately.

# Warnings and Cautions

## Avoid explosive or flammable gases

It has been determined that the SAM 360P is safe to use with oxygen mask delivery systems. However, to avoid the risk of an explosion, it is strongly advised that you do NOT use the SAM 360P in the vicinity of explosive gases, including flammable anaesthetics or concentrated oxygen.



## Caution

### Correct placement of the electrode pads

Proper placement of the SAM 360P electrode pads is critical. You must strictly observe the instructions shown in the Emergency User Guide and on the device. Wrong placement, or the presence of air, hair, surgical dressings or medicine patches between the pads and the skin, could reduce defibrillation effectiveness. Slightly red skin after shock therapy is normal.

### Do not touch the patient during analysis

Touching the patient during the analysis phase of treatment can cause interference with the diagnostic process. Avoid contact with the patient while analysis is being carried out. The device will instruct you when it is safe to touch the patient.

## **Do not use if the pouch containing the electrodes is not sealed**

The Pad-Pak is a single-use item and you must replace it after each use or if the pouch that seals defibrillation pads has been broken or compromised in any way. If you suspect that the Pad-Pak is damaged, you must replace it immediately.



## **Notice**

### **Susceptibility to electromagnetic interference**

To safeguard against interference, you must operate the SAM 360P at least 2 m (6 feet) away from all radio frequency devices. Alternatively, switch off the equipment causing the electromagnetic interference.

### **Temperature range for operation**

The SAM 360P, with its battery, pads and electrodes, is designed to operate in the temperature range of 0 °C to 50 °C (50 °F to 122 °F). Use of the device outside this range may cause malfunction.

### **Ingress protection**

The IP56 rating does not cover the immersion of any part of the SAM 360P in water or any type of fluid. Contact with fluids may seriously damage the device or cause fire or a shock hazard.

# Warnings and Cautions

## Prolonging battery life

Do not turn on the device unnecessarily as this may reduce the standby life of the device.

Standby storage outside the range of 0 °C to 50 °C (50 °F to 122 °F) may decrease the shelf-life of the Pad-Pak.

## Do not test on simulators and manikins

Our devices cannot be tested using industry-standard simulators and manikins.

Our algorithm uses heart rate variability as one of its criteria for measuring ventricular fibrillation (VF). Consequently we do not recommend the use of normal simulators to test our device.



## Further Information

### Use of this manual

It is important that you read this manual carefully before using the SAM 360P. This manual is presented in support of any training you may have received. If you have any questions, contact your authorised distributor or HeartSine Technologies directly for advice or explanation.

The information in this manual is subject to change without notice and does not represent a commitment on behalf of HeartSine Technologies. No part of this manual may be reproduced or transmitted in any form or by any means, electrical or mechanical, including photocopying and recording, for any purpose without the express written permission of HeartSine Technologies.



## **Operator training**

The SAM 360P is intended for use by personnel who have been trained in its operation. Users should have received training in basic life support / AED, advanced life support or a physician-authorized emergency medical response training program.

## **Use of accessories**

The SAM 360P is a self-contained device. Do not use any unauthorized accessories with it. The SAM 360P may malfunction if non-approved accessories are used.

## **Regular maintenance**

Check the device periodically. See 'Service and Maintenance' on page 21.

## **Correct disposal of the device**

Dispose of the device in accordance with your national or local regulations, or contact your HeartSine distributor. Please follow the 'After use' on page 17.

## **Compliance with local regulations**

Check with the relevant local government health department for information about any requirements associated with ownership and use of a defibrillator in the region where it is to be used.

# Introduction

## The SAM 360P

The SAM 360P is a fully-automatic (there is no shock button to press), external defibrillator designed to quickly deliver a defibrillation shock to victims of sudden cardiac arrest (SCA).

The SAM 360P is designed to operate in accordance with the joint European Resuscitation Council (ERC) and American Heart Association (AHA) 2015 guidelines on Cardiopulmonary Resuscitation (CPR) and Emergency Cardiovascular Care (ECC).

## Sudden cardiac arrest (SCA)

Sudden cardiac arrest is a condition in which the heart suddenly stops pumping effectively due to a malfunction of the heart's electrical system. Often victims of SCA have no prior warning signs or symptoms. SCA can also occur in people with previously diagnosed heart conditions. Survival from SCA depends on immediate and effective cardiopulmonary resuscitation (CPR).

The use of an external defibrillator within the first few minutes of collapse can greatly improve patient's chances of survival. Heart attack and SCA are not the same, though sometimes a heart attack can lead to an SCA. If you are experiencing symptoms of a heart attack (chest pain, pressure, shortness of breath, tight feeling in the chest or elsewhere in the body), seek emergency medical attention immediately.

## Ventricular fibrillation

The normal electrical rhythm by which the heart muscle contracts to create blood flow around the body is known as normal sinus rhythm (NSR). Ventricular fibrillation (VF), caused by chaotic electrical signals in the heart, is often the cause of SCA. In victims of SCA it is possible to re-establish normal sinus rhythm by means of an electric shock across the heart. This treatment is called defibrillation.

## Recommended training

SCA is a condition requiring immediate emergency medical intervention. Due to the nature of the condition, this intervention can be performed before seeking the advice of a physician.

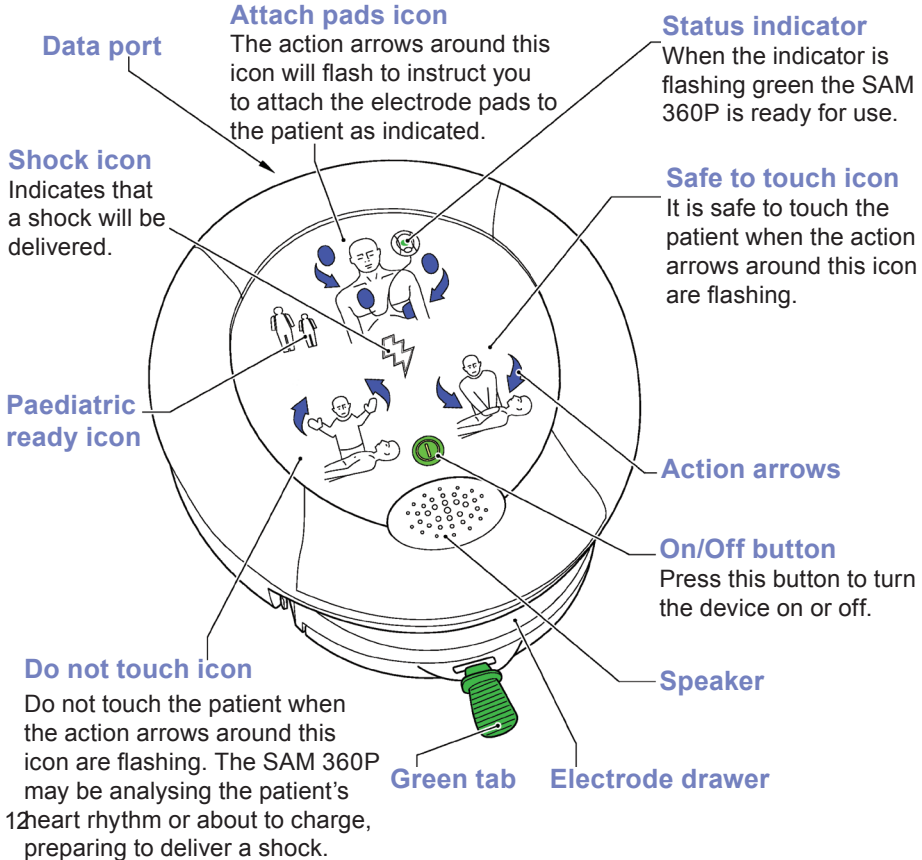
To properly diagnose this condition, HeartSine recommends that all potential users of the SAM 360P are fully trained in cardiopulmonary resuscitation (CPR), basic life support (BLS) and, in particular, the use of an automated external defibrillator. HeartSine also recommends that this training be kept up to date by regular refresher courses as and when recommended by your training provider.

If potential users of the SAM 360P are not trained in these techniques, contact your authorised distributor or HeartSine Technologies directly. Either can arrange for training to be provided. Alternatively contact your local government health department for information on certified training organisations in your region.

## CPR metronome

During CPR the SAM 360P will play an audible beep and flash the “Safe To Touch” indicator at a rate compliant with 2015 AHA/ERC guidelines. This feature is referred to as the CPR metronome. Use the metronome as a guide on how frequently to compress a patient’s chest if you need to apply CPR.

# SAM 360P Overview

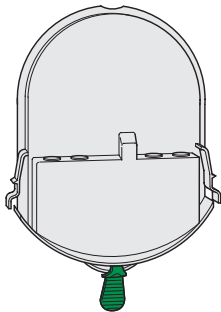


# Preparation

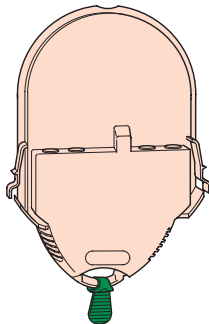
## Unpacking

Check that the contents include the User Manual, soft case, Pad-Pak, Warranty Card and Emergency User Guide.

A Pad-Pak is a single-use removable battery and electrode pack in one unit. It is available in two versions<sup>1</sup>: grey coloured Pad-Pak for use with adults and a pink coloured Paediatric-Pak for use with children (see the illustration below).



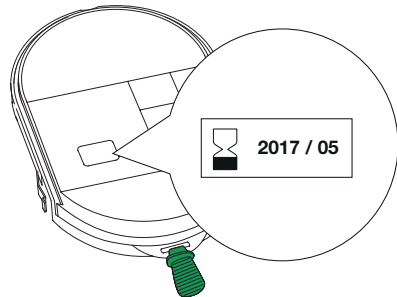
Adult Pad-Pak



Paediatric-Pak

## Checks before putting into service

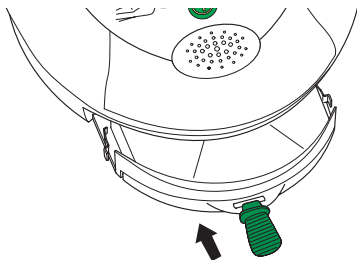
1. Check the expiry date (year/month) on the rear of the Pad-Pak (see the illustration below). If the expiry date has passed, you must replace the Pad-Pak.




<sup>1</sup> A third version is also available specifically for airlines

# Preparation

- Unpack the Pad-Pak. Retain the packaging in case you need to return the Pad-Pak to HeartSine. Place the SAM 360P on a flat surface. Insert the Pad-Pak into the SAM 360P (see the illustration below). Listen for the “click” sound and ensure both tabs are fully engaged.




- If required, the SAM 360P will run a self test routine. The action arrows will flash during this process. On successful completion of the self-test routine, the green status indicator (see ‘SAM 360P Overview’ on page 12) will blink. If so, your SAM 360P is ready for use.

- Turn on the SAM 360P by pressing  on the front panel to check that the device is operating correctly. Listen for the voice prompts but do **NOT** follow them. Make sure that no warning messages are played.



**Notice: Do NOT pull the green tab on the Pad-Pak. If you have opened the electrode drawer, you may have to replace your Pad-Pak.**

**Only turn the SAM 360P on ONCE. If you turn it on and off repeatedly, you will exhaust the batteries prematurely and you may need to replace the Pad-Pak.**

- Turn off the SAM 360P by pressing  on the front panel. Check the status indicator (see ‘SAM 360P Overview’ on page 12) is flashing green. If you have heard no warning messages and the status indicator is flashing green, the device is ready for use.

6. Place the SAM 360P in its supplied soft carry case. Store the SAM 360P in an unobstructed, secure location in a **clean, dry environment** specifically where it will be seen and heard. Be sure to store according to specifications (see 'Technical Data' on page 26).

Standby temperature: 0 °C to 50 °C  
(50 °F to 122 °F)

Relative humidity: 5% to 95%  
(non-condensing)



**Notice: HeartSine recommends that you keep a spare Pad-Pak with your SAM 360P. You can store it in the rear section of the soft carry case.**

7. Complete the Warranty Card and return it to your authorised distributor or HeartSine Technologies directly (see 'Tracking Requirements' on page 22).

## Preparation checklist

- Step 1. Check the Pad-Pak expiry date.
- Step 2. Install the Pad-Pak.
- Step 3. Check for a successful completion of the self-test routine.
- Step 4. Turn on to check operation.
- Step 5. Turn off.
- Step 6. Store the SAM 360P correctly.
- Step 7. Register your SAM 360P.
- Step 8. Create a service schedule (see 'Service and Maintenance' on page 21).

# Using the SAM 360P

## When to use

The SAM 360P is indicated for use on victims of sudden cardiac arrest who are exhibiting the following signs:

- Unconscious
- Not breathing
- Without circulation

The SAM 360P has been designed to work on unconscious, nonresponsive patients. If the patient is responsive or conscious, do not use the SAM 360P to provide treatment.

The SAM 360P is suitable for use on patients of over 25 kg (55 lbs) in weight or equivalent to a child of approximately eight years old or over.

For use on smaller children (from 1 to 8 years old), remove the adult Pad-Pak and install a Paediatric-Pak.

If a Paediatric-Pak or an alternative suitable defibrillator is not available, you may use an adult Pad-Pak.

## Using the SAM 360P


Refer to the separate Emergency User Guide. During use the SAM 360P will give extensive voice prompts to guide a user. For full list of voice prompts see 'List of Voice Prompts' on page 38.

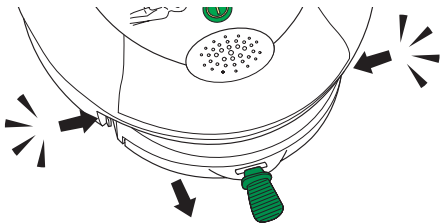


**Notice: The SAM 360P aborts a ready to shock condition once a non-shockable rhythm is detected.**



## After use

1. Turn off the SAM 360P by pressing  on the front panel.
2. Remove the electrode pads from the patient and stick them together 'face to face'. The electrodes may be contaminated with human bodily tissue, fluid or blood. Dispose of the electrodes separately as an infectious waste material.
3. The Pad-Pak contains Lithium batteries. It is a single-use item and must be replaced after each use. Remove the Pad-Pak by pressing the two tabs on either side of the Pad-Pak. The Pad-Pak will slide forward (see the illustration below).



Do not dispose of the SAM 360P or Pad-Pak in the normal waste. Dispose of it at an appropriate recycling facility according to local requirements. Alternatively return it to your distributor for disposal or replacement.

4. Check the SAM 360P for dirt or contamination. If necessary, clean it using a soft cloth dampened by one of the following:

Soapy water

Isopropyl alcohol (70% solution).



**Caution: Do not immerse any part of the SAM 360P in water or any type of fluid. Contact with fluids may seriously damage the device or cause a fire or a shock hazard.**



**Notice: Do not clean the SAM 360P with abrasive materials, cleaners or solvents.**

# Using the SAM 360P

5. Check the SAM 360P for damage. If the SAM 360P is damaged, replace it immediately.
6. Install a new Pad-Pak. Before installing, check the Pad-Pak expiry date (see 'Preparation' on page 13). After installation check the status indicator is blinking green.

# Paediatric-Pak

## Using the Paediatric-Pak

The Paediatric-Pak is intended to provide therapy for paediatric (child) victims of SCA between the ages of 1 and 8 years old who are:

Unconscious

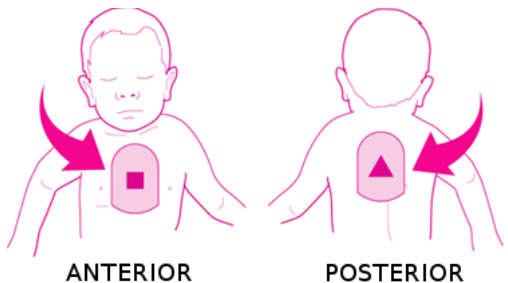
Not breathing

Without circulation

## Electrode Placement:

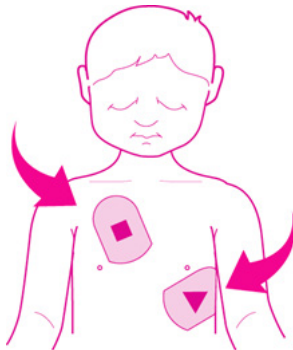
For paediatric patients there are two options for electrode placement:

a. If a child's chest is small it may be necessary to place one pad on the child's BARE chest in the center, and the other pad on the child's BARE back in the center of the ribcage as shown in Method a).



Method a)

b. If a child's chest is large enough to permit a 2.5cm (1 inch) gap between the pads, pad placement can be used similar to adult placement. Place one pad on child's BARE upper right chest above nipple and one pad on child's BARE lower left ribcage below nipple as shown in Method b).



Method b)

Electrodes can be placed on the child's chest if their chest is large enough OR if trauma does not allow for the placement as shown in Method a).



**Warning: Defibrillation electrodes must be at least approx 2.5 cm (1 inch) apart and should never be touching one another.**



**Warning: The Paediatric-Pak contains a magnetic component (surface strength 6500 gauss). Avoid storage next to magnetically sensitive storage media.**



**Warning: Not for use on patients under 1 year old. For use with children up to the age of 8 years or up to 25kg (55lbs). DO NOT DELAY THERAPY IF YOU ARE NOT SURE OF EXACT AGE OR WEIGHT.**

# Service and Maintenance

HeartSine recommends users perform regular maintenance checks. The recommended maintenance checks are:

## **Weekly**

- Check the status indicator. If the green status indicator is not flashing every 5 to 10 seconds or if the red status indicator is flashing or if you hear beeping, a problem has been detected. See 'Troubleshooting' on page 24. The SAM 360P performs a self-test routine at midnight GMT every Sunday. During this self-test the status light blinks red but returns to green on successful completion of the self-test routine. The self-test takes no more than 10 seconds to complete. If the status indicator continues to flash red the SAM 360P has a fault (see 'Troubleshooting' on page 24).

## **Monthly**

- If the device shows any signs of physical damage, contact your authorised distributor or HeartSine Technologies directly.
- Check the expiry date of the SAM 360P Pad-Pak (see 'Preparation' on page 13 for the location of the date). If the date has expired, or is near expiry, replace with a new Pad-Pak or contact your local HeartSine distributor for a replacement.

If you hear a warning message when you turn on your SAM 360P or if, for any reason, you have suspicions that your SAM 360P is not working correctly, read the section 'Troubleshooting' on page 24.

# Tracking Requirements

Medical Devices Regulations require us to track the location of all medical devices sold.

It is important that you complete the samaritan PAD Warranty Card with your details and return it to your authorised distributor or HeartSine Technologies directly.

Alternatively send an email, to [support@heartsine.com](mailto:support@heartsine.com), containing:

Name

Address

Device serial number

or use our on-line registration tool at <https://secure.heartsine.com/UserRegistration.html>

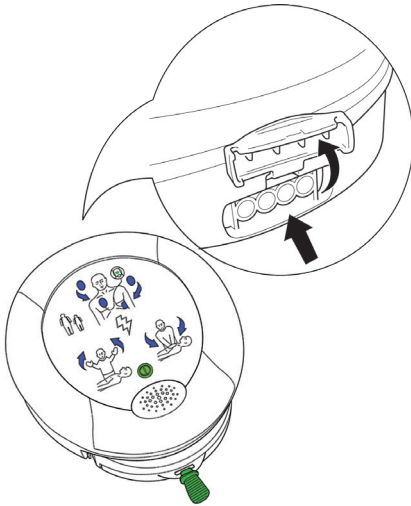
Your participation will allow us to contact you with any important notifications about the SAM 360P, such as any future software updates or field safety corrective actions.

If there is a change in the information you have provided to us, such as a change of address or a change in ownership of your SAM 360P, contact us with the updated information.

# Data Management

The HeartSine Saver EVO™ software is an optional accessory. Contact your authorised distributor or HeartSine Technologies directly about the after-use data management service.

1. Connect the USB cable to the SAM 360P (see illustration below).



2. Connect the USB cable to a PC.
3. Launch the HeartSine Saver EVO utility.



**Notice: The SAM 360P should only be connected to an IEC60950 PC.**





**Caution: You cannot defibrillate while the SAM 360P is connected to a PC.**

For further information on this optional accessory, contact your authorised distributor or HeartSine Technologies directly.

# Troubleshooting

## Status indicator flashing red

If the status indicator is flashing red or if the device is emitting a 'beep', check the expiry date on your Pad-Pak (see 'Preparation' on page 13). If the expiry date has not been passed, turn on the SAM 360P by pressing  on the front panel and listen for the voice prompt 'call for medical assistance'. Then turn off by pressing  on the front panel. If this action does not correct the problem, contact your authorised distributor or HeartSine Technologies immediately.

## Low battery warning



**This message does not indicate a fault.**

The first time the device plays the message 'warning low battery', it will still continue to function properly. However, it may have fewer than 10 shocks left. If you hear this message, prepare the spare Pad-Pak for use and be prepared to swap it quickly. Order a new Pad-Pak as soon as possible.

## Memory full warning

If the device plays the message 'memory full', then the memory can record no further ECG data or events. However, the device can still analyse and deliver a shock if required. If you hear this message, contact HeartSine Technologies technical support.

## Audible warnings

If the device emits 3 beeps rapidly when turned off, it has sensed that the ambient temperature is outside of the specified operating range. This beeping could also occur during the weekly self-test. If you hear this beeping, please ensure the device is returned to the specified operating conditions.

During use, if the status indicator changes from green to red and the device starts to 'beep', there is insufficient battery capacity to deliver a shock. The device will continue to analyse the patient's heart rhythm and advise when CPR is needed.



## Device service required

If the device plays the message 'device service required', then it has detected a fault. Contact your authorised distributor or HeartSine directly for further instruction.



**Warning: If you hear this message during use, seek an alternative defibrillator immediately.**

**No modification of this equipment is allowed.**

## Sources of support

If you have completed the troubleshooting steps above and you find the device is still not working correctly, contact your authorised distributor or HeartSine Technologies Technical Support at [support@HeartSine.com](mailto:support@HeartSine.com).

## Warranty exclusion

HeartSine or its authorised distributors are not obliged to replace or repair under warranty if one or more of the following conditions apply:

The device has been opened.

Unauthorised modifications have been made.

The device has not been used in accordance with the instructions provided in this manual.

The serial number has been removed, defaced, altered or, by any other means, made unreadable.

The device has been used or stored outside its indicated temperature range.

The Pad-Pak packaging is not returned.

The device has been tested using unapproved methods or inappropriate equipment, (see 'Warnings and Cautions' on page 5).

# Technical Data

## Physical parameters (with Pad-Pak installed)

Size: 20 cm x 18.4 cm x 4.8 cm (8.0 in x 7.25 in x 1.9 in)  
Weight: 1.1 kg (2.4 lbs)

## Environmental

Operating temperature: 0 °C to 50 °C (32 °F to 122 °F)  
Standby temperature: 0 °C to 50 °C (32 °F to 122 °F)  
Transport temperature: -10 °C to 50 °C (14 °F to 122 °F) for up to two days. If the device has been stored below 0 °C (32 °F), it should be returned to an ambient temperature of between 0 °C to 50 °C (32 °F to 122 °F) for at least 24 hours before use.  
Relative humidity: 5% to 95% (non-condensing)  
Enclosure: IEC 60529/EN 60529 IP56  
Altitude: 0 metres to 4575 metres (0 feet to 15 000 feet)  
Shock: MIL STD 810F Method 516.5, Procedure 1 (40G's)  
Vibration: MIL STD 810F Method 514.5 Procedure 1 Category 4  
MIL STD 810F Method 514.5 Procedure 1 Category 7

## Pad-Pak and Paediatric-Pak

|                             |   |
|-----------------------------|---|
| Weight:                     | 0.2 kg (0.44 lbs)   |
| Battery type:               | Disposable single-use combined battery and defibrillation electrode cartridge (lithium manganese dioxide (LiMnO <sub>2</sub> ) 18V) |
| Battery capacity (new):     | >60 shocks or 6 hours of continuous monitoring  |
| Battery capacity (4 years): | >10 shocks  |
| Standby life:               | See the expiry date on the Pad-Pak.   |
| Electrode type:             | Single-use pre-attached combined ECG sensor/defibrillation pad  |
| Electrode placement:        | Adult: anterior-lateral<br>Paediatric: electrodes anterior-posterior or anterior-lateral  |
| Electrode active area:      | 100 cm <sup>2</sup> (40 inch <sup>2</sup> )   |
| Electrode cable length:     | 1 m (3.25 ft)   |
| Electrode shelf life:       | See the expiry date on the Pad-Pak.   |

## Patient analysis system

|                          |   |
|--------------------------|---|
| Method:                  | Evaluates the patient's ECG, signal quality, electrode contact integrity and patient impedance to determine if defibrillation is required |
| Sensitivity/Specificity: | Meets IEC 60601-2-4   |

# Technical Data

## User Interface

|                  |   |
|------------------|---|
| Visual prompts:  | Attach pads, stand clear, perform CPR, shock now, self test pass - ready state                                  |
| Audible prompts: | Extensive voice prompts guide the user through the operation sequence (see 'List of Voice Prompts' on page 38). |
| Languages:       | Contact your HeartSine authorised distributor.  |
| Controls:        | One button: 'On/Off'  |

## Defibrillator performance

|  |   |
|--|---|
| Times to shock delivery (fresh battery or after 6 shocks): |   |
| Charging time:   | Typically 150J in < 8 sec, 200J in < 12 sec |
| Following CPR:   | Typically 19 seconds                        |
| Impedance range:   | 20 $\Omega$ to 230 $\Omega$                 |

## Therapeutic shock

|           |  |
|-----------|--|
| Waveform: | SCOPE™ (Self Compensating Output Pulse Envelope) biphasic escalating waveform. Optimised biphasic waveform compensates energy, slope and envelope for patient impedance                      |
| Energy:   | Pre-configured factory settings for escalating energy are Version AHA/ERC 2015<br>Adult: Shock 1: 150J; Shock 2: 150J; Shock 3: 200J<br>Paediatric: Shock 1: 50J; Shock 2: 50J; Shock 3: 50J |

## Event recording

|         |  |
|---------|--|
| Type:   | Internal memory  |
| Memory: | 90 minutes of ECG (full disclosure) and event/incident recording                             |
| Review: | Custom USB cable directly connected to a PC and Saver EVO Windows-based data review software |

## Electromagnetic compatibility

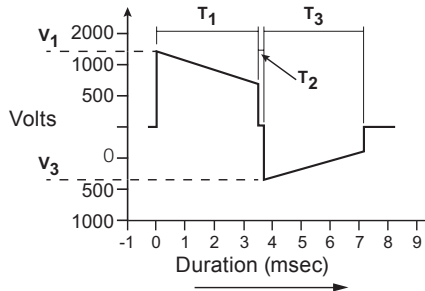
|                          |   |
|--------------------------|---|
| EMC:                     | IEC60601-1-2  |
| Radiated Emissions:      | IEC55011  |
| Electrostatic Discharge: | IEC61000-4-2 (8 kV)   |
| RF Immunity:             | IEC61000-4-3 80 MHz – 2.5 GHz, (10 V/m)                           |
| Magnetic Field Immunity: | IEC61000-4-8 (3 A/m)  |
| Aircraft:                | RTCA/DO-160F, Section 21 (Category M)<br>RTCA DO-227 (ETSO-C142a) |

# Technical Data

## SCOPE™ Biphasic Waveform

The SAM 360P delivers a Self Compensating Output Pulse Envelope (SCOPE) biphasic waveform. This waveform automatically optimises the waveform pulse envelope (amplitude, slope, and duration) for a wide range of patient impedances, from 20 ohms to 230 ohms. The delivered waveform to the patient is an optimised, impedance-compensated, biphasic, truncated exponential waveform that incorporates an escalating energy protocol of 150 joules, 150 joules, and 200 joules. The duration of each phase is automatically adjusted to compensate for varying patient impedances. The first phase (T1) duration is always equivalent to the second phase (T3) duration. The interphase pause (T2) is always a constant 0.4 ms for all patient impedances.

The specific SCOPE waveform characteristics for a 150 joules pulse are listed opposite.



| Resistance (ohms) | Waveform Voltages (volts) |        | Waveform Duration (ms) |                |
|-------------------|---------------------------|--------|------------------------|----------------|
|                   | V <sub>1</sub>            | Tilt % | T <sub>1</sub>         | T <sub>3</sub> |
| 25                | 1640                      | 63.1   | 3                      | 3              |
| 50                | 1650                      | 52.7   | 4.5                    | 4.5            |
| 75                | 1660                      | 51.4   | 6.5                    | 6.5            |
| 100               | 1670                      | 48.7   | 8                      | 8              |
| 125               | 1670                      | 50.4   | 10.5                   | 10.5           |
| 150               | 1670                      | 48.7   | 12                     | 12             |
| 175               | 1670                      | 48.7   | 14                     | 14             |
| 200               | 1670                      | 47.6   | 15.5                   | 15.5           |
| 225               | 1680                      | 46.7   | 17                     | 17             |

Adult Pad-Pak waveform specification

All values are nominal

| Resistance<br>(ohms) | Energy<br>(joules) | Waveform<br>Voltages<br>(volts) |           | Waveform<br>Duration<br>(ms) |                |
|----------------------|--------------------|---------------------------------|-----------|------------------------------|----------------|
|                      |                    | V <sub>1</sub>                  | Tilt<br>% | T <sub>1</sub>               | T <sub>3</sub> |
| 25                   | 47.5               | 514                             | 55.6      | 7.8                          | 5.4            |
| 50                   | 51.3               | 671                             | 50.4      | 8.8                          | 6              |
| 75                   | 52.1               | 751                             | 47.1      | 10                           | 6.6            |
| 100                  | 51.8               | 813                             | 44.3      | 10.8                         | 6.8            |
| 125                  | 52.4               | 858                             | 41.4      | 11.5                         | 7.3            |

Paediatric-Pak waveform specification

All values are nominal

### Motion detection algorithm \*

The SAM 360P uses the HeartSine samaritan ICG analysis to detect chest compression artefact and other forms of motion that prompts an audio warning and instruction to stop CPR or other motion.

\* *Motion detection algorithm performance may be reduced during low battery operation.*

### Arrhythmia analysis algorithm

The SAM 360P uses the HeartSine samaritan ECG arrhythmia analysis algorithm. This algorithm will evaluate the patient's ECG to ascertain if a therapeutic shock is appropriate. If a shock is required, the SAM 360P will charge and advise the user to stand clear. If no shock is advised, the device will pause to allow the user to deliver CPR.

The SAM 360P ECG arrhythmia analysis algorithm performance has been extensively evaluated by using several databases of real-life ECG traces. Included in this are the American Heart Association's (AHA) database and the Massachusetts Institute of Technology MIT – NST database. The SAM 360P ECG arrhythmia analysis algorithm's sensitivity and specificity meet the requirements of IEC60601-2-4.

# Technical Data

The SAM 360P ECG arrhythmia analysis algorithm performance is summarised in the table below:

| Rhythm Class  | ECG Test Sample Size (seconds) | Required Performance Specifications | Performance Results (%) | 90% One-Sided Lower Confidence Limit |
|---|--------------------------------|-------------------------------------|-------------------------|--------------------------------------|
| Shockable Rhythm:<br>Ventricular Fibrillation (VF)      | 14538                          | Sensitivity > 90%                   | 97.03                   | 95.31                                |
| Shockable Rhythm:<br>Ventricular Tachycardia (VT)       | 2321                           | Sensitivity > 75%                   | 92.55                   | 80.52                                |
| Non-Shockable Rhythm:<br>Combined Non-Shockable Rhythms | 301177                         | Specificity > 95%                   | 98.30                   | 95.02                                |



### Guidance and manufacturer's declaration – electromagnetic emissions

The SAM 360P is intended for use in the electromagnetic environment specified below. The customer or user of the SAM 360P must assure that it is used in such an environment.

| Emissions test  | Compliance     | Electromagnetic environment – guidance  |
|---|----------------|---|
| RF emissions CISPR 11                                   | Group 1        | The SAM 360P uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.                       |
| RF emissions CISPR 11                                   | Class B        | The device is suitable for use in all establishments, including domestic and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes |
| Harmonic emissions IEC/EN 61000-3-2                     | Not applicable |   |
| Voltage fluctuations/flicker emissions IEC/EN 61000-3-3 | Not applicable |   |

# Technical Data

## Guidance and manufacturer's declaration – electromagnetic immunity

The SAM 360P is intended for use in the electromagnetic environment specified below. The customer or user of the SAM 360P must assure that it is used in such an environment.

| Immunity test   | IEC 60601 test level   | Compliance level             | Electromagnetic environment – guidance  |
|---|--|------------------------------|---|
| Electrostatic discharge (ESD)<br>IEC/EN 61000-4-2   | ± 6 kV contact<br>± 8 kV air   | ± 6 kV contact<br>± 8 kV air | Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%. |
| Electrical fast transient/burst<br>IEC/EN 61000-4-4   | ±2kV for power supply lines<br>±1 kV for input/output lines  | Not Applicable               | Not Applicable  |
| Surge<br>IEC/EN 61000-4-5   | ±1kV differential mode<br>±2 kV common mode  | Not Applicable               | Not Applicable  |
| Voltage dips, short interruptions and voltage variations on power supply input lines<br>IEC/EN 61000-4-11 | <5 % Ut (>95 % dip in Ut) for 0.5 cycle<br>40 % Ut (60% dip in Ut) for 5 cycles<br>70 % Ut (30 % dip in Ut) for 25 cycles<br><5 % Ut (>95 % dip in Ut) for 5 sec | Not Applicable               | Not Applicable  |
| Power-frequency (50/60 Hz) magnetic field<br>IEC/EN 61000-4-8   | 3 A/m  | 3A/m                         | Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.     |

Note: Ut is the a.c.mains voltage prior to application of the test level

## Guidance and manufacturer's declaration – electromagnetic immunity

The SAM 360P is intended for use in the electromagnetic environment specified below. The customer or user of the SAM 360P must assure that it is used in such an environment.

| Immunity test                    | IEC 60601 test level  | Compliance level               | Electromagnetic environment – guidance   |
|----------------------------------|---|--------------------------------|--|
| Conducted RF<br>IEC/EN 61000-4-6 | 3 Vrms<br>150 kHz to 80 MHz<br>outside ISM bands <sup>a</sup> | Not applicable                 | <p>Portable and mobile RF communications equipment should be used no closer to any part of the SAM 360P, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.</p> <p><b>Recommended separation distance</b></p> <p>Not applicable</p>   |
|                                  | 10 Vrms<br>150 kHz to 80 MHz<br>in ISM bands <sup>a</sup>     | Not applicable                 | Not applicable   |
| Radiated RF<br>IEC/EN 61000-4-3  | 10 V/m<br>80 MHz to 2.5 GHz                                   | 10 V/m<br>80 MHz<br>to 2.5 GHz | <p><math>d = 1.2 \sqrt{P}</math> 80 MHz to 800 MHz<br/> <math>d = 2.3 \sqrt{P}</math> 800 MHz to 2.5 GHz</p> <p>Where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in metres (m)<sup>b</sup>.</p> <p>Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey<sup>c</sup>, should be less than the ... [next page]</p> |

# Technical Data

## Guidance and manufacturer's declaration – electromagnetic immunity

... compliance level in each frequency range.  
Interference may occur in the vicinity of equipment marked with the following symbol:



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

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

a The ISM (industrial, scientific and medical) bands between 150kHz and 80 MHz are 6,765 MHz to 6,795 MHz; 13,553 MHz to 13,567 MHz; 26,957 MHz to 27,283 MHz; 40,66 MHz to 40,70 MHz;

b The compliance levels in the ISM frequency bands between 150 kHz and 80 MHz and in the frequency range 80 MHz to 2.5 GHz are intended to decrease the likelihood that mobile/portable communications equipment could cause interference if it is inadvertently brought into patient areas. For this reason, an additional factor of 10/3 has been incorporated into the formulae used in calculating the recommended separation distance for transmitters in these frequency ranges.

c Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location where the SAM 360P is used exceeds the applicable RF compliance level (see above), the SAM 360P should be observed to verify that it is operating normally. If abnormal operation is observed, additional measures may be necessary, such as re-orienting or relocating the SAM 360P.

**Recommended separation distances between portable and mobile RF communication equipment and the SAM 360P**

The SAM 360P is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the SAM 360P can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the SAM 360P as recommended below, according to the maximum output power of the communications equipment.

| Rated maximum output power of transmitter<br><b>W</b> | Separation distance according to frequency of transmitter<br><b>m</b> |                                   |   |   |
|---|---|-----------------------------------|---|---|
|   | 150 kHz to 80 MHz<br>outside ISM bands                                | 150 kHz to 80 MHz<br>in ISM bands | 80 MHz to 800 MHz<br>$d = 1.2 \sqrt{P}$ | 800 MHz to 2.5GHz<br>$d = 2.3 \sqrt{P}$ |
| 0.01  | Not applicable  | Not applicable                    | 0.12                                    | 0.23                                    |
| 0.1   | Not applicable  | Not applicable                    | 0.38                                    | 0.73                                    |
| 1   | Not Applicable  | Not Applicable                    | 1.2                                     | 2.3                                     |
| 10  | Not Applicable  | Not Applicable                    | 3.8                                     | 7.3                                     |
| 100   | Not Applicable  | Not Applicable                    | 12                                      | 23                                      |

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

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

NOTE 2 The ISM (industrial, scientific and medical) bands between 150kHz and 80 MHz are 6,765 MHz to 6,795 MHz; 13,553 MHz to 13,567 MHz; 26,957 MHz to 27,283 MHz; 40,66 MHz to 40,70 MHz.

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

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

# List of Voice Prompts

Listed below are the voice prompts used by the SAM 360P. Read the voice prompts in advance of use to be familiar with the types of instructions given.

## Adult patient/child patient

- “Call for medical assistance”
- “Remove clothing from patient’s chest to expose bare skin”
- “Pull ‘green tab’ to remove pads”
- “Peel pads from liner”
- “Apply pads to patient’s bare chest as shown in picture”
- “Press pads firmly to patient’s bare skin”
- “Assessing heart rhythm – do not touch the patient”
- “Analysing - do not touch the patient”

## If motion is detected...

- “Motion detected – do not touch the patient”

## If a shock is not required...

- “No shock advised”
- “Begin CPR”
- “It is safe to touch the patient”
- “Place overlapping hands in middle of chest”
- “Press directly down on the chest in time with the metronome”
- “Remain calm”

## If a shock is required...

- “Stand clear of patient – shock advised”
- “Stand clear of patient – shock will be delivered in 3,2,1”
- “Shock delivered”
- “Begin CPR”
- “It is safe to touch the patient”
- “Place overlapping hands in middle of chest”
- “Press directly down on the chest in time with the metronome”
- “Remain calm”

# Notes

Authorised Distributor

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