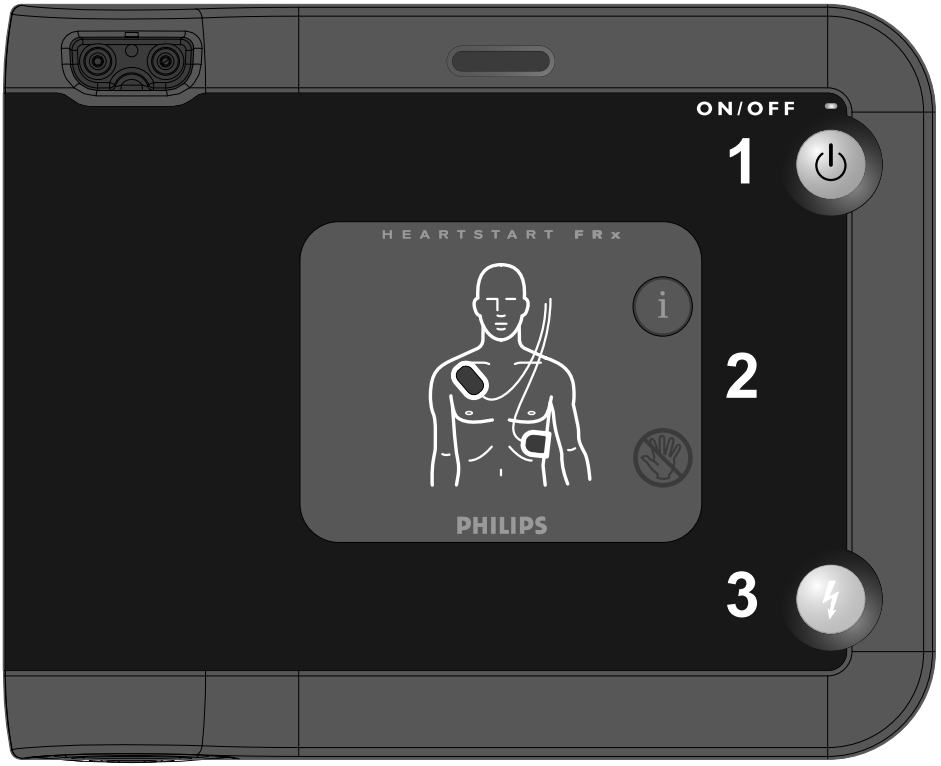
HEARTSTART FRx DEFIBRILLATOR

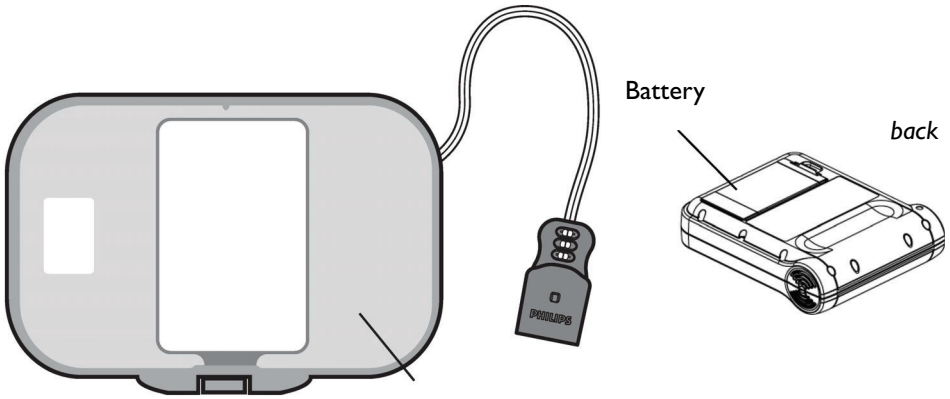
OWNER'S MANUAL



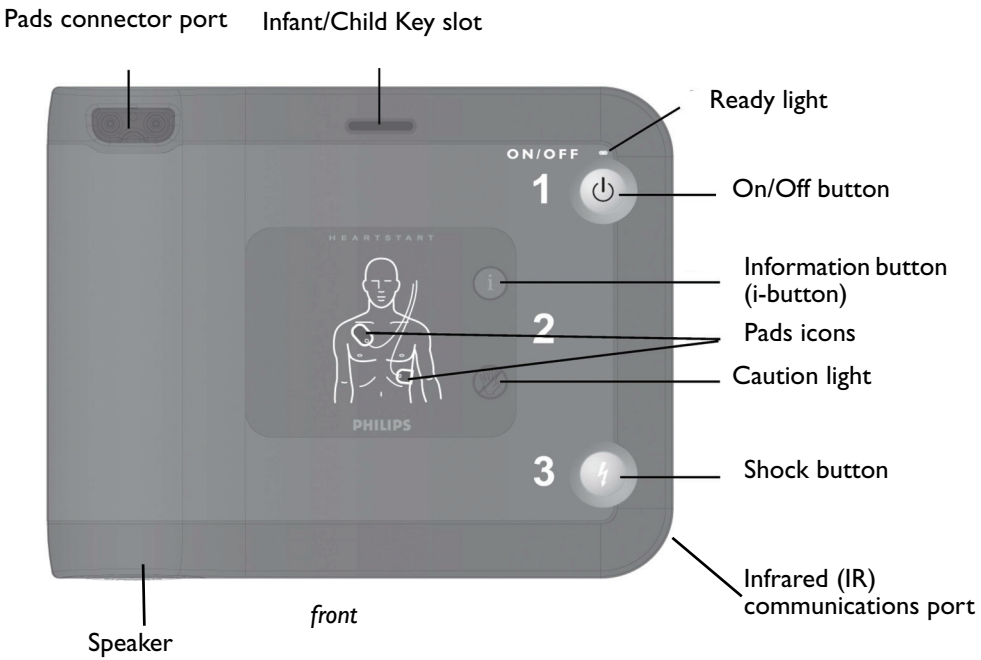
REF 861304
Edition 15

PHILIPS

Intentionally blank.



SMART PADS II in closed case



The HeartStart FRx Defibrillator 861304

Intentionally blank.

HeartStart FRx Defibrillator
QUICK REFERENCE



PHILIPS MEDICAL SYSTEMS

PHILIPS

Intentionally blank.

HeartStart FRx

REF 861304

Automated External Defibrillator

OWNER'S MANUAL

Edition 15

IMPORTANT NOTE:

It is important to understand that survival rates for sudden cardiac arrest are directly related to how soon victims receive treatment. For every minute of delay, the chance of survival declines by 7% to 10%.

Treatment cannot assure survival. In some victims, the underlying problem causing the cardiac arrest is simply not survivable despite any available care.

PHILIPS MEDICAL SYSTEMS

PHILIPS

Intentionally blank.

About This Edition

The information in this guide applies to the HeartStart FRx Defibrillator 861304. This information is subject to change. Please contact Philips at www.philips.com/AEDsupport or your local Philips representative for information on revisions.

Edition History

Edition 15
Publication Date: March 2023
Publication Number: 45356498251 1

Notices

© 2023 Koninklijke Philips N.V.
All rights reserved.

Specifications are subject to change without notice.

Trademarks are the property of Koninklijke Philips N.V. or their respective owners.

CAUTION: Federal law (USA) restricts this device to sale by or on the order of a physician.

Rx only

The Philips HeartStart FRx is designed to be used only with Philips-approved accessories. The FRx may perform improperly if non-approved accessories are used.

Device Tracking

In the USA, this device is subject to tracking requirements by the manufacturer and distributors. If the defibrillator has been sold, donated, lost, stolen, exported, or destroyed, notify Philips Medical Systems or your distributor.

Device Manufacturer

Philips Medical Systems,
22100 Bothell Everett Highway
Bothell, WA, 98021-8431, USA

Patents

Patents listing at www.ip.philips.com/patentmarking.

For Technical Support

If you need technical support, please contact your local Philips representative by calling the regional number on the back cover of this manual, or go to www.philips.com/AEDsupport.

To download additional copies of this manual, go to www.philips.com/IFU.

PHILIPS

PHILIPS MEDICAL SYSTEMS

Intentionally blank.

CONTENTS

1 INTRODUCTION TO THE HEARTSTART FRX

Description 1
Sudden cardiac arrest 1
Indications for use 2
Contraindications 2
Dangers, Warnings, and Cautions 2
Probable Adverse Effects of the Device on Health 7
Clinical Summary of Safety & Performance Data 7
Pediatric Defibrillation Waveform 11
Essential Performance 13
Principles of Operation 13
Implementation considerations 14
For more information 14

2 SETTING UP THE HEARTSTART FRX

Package contents 15
Setting up the FRx 15
Recommended accessories 18

3 USING THE HEARTSTART FRX

Overview 19
STEP 1: Press the green on/off button 20
STEP 2: Follow the FRx voice instructions 20
STEP 3: Press the flashing orange Shock button if instructed 21
Treating infants and children 22
When emergency medical services arrive 24

4 AFTER USING THE HEARTSTART FRX

After each use 27
FRx data storage 28

5 MAINTAINING THE HEARTSTART FRX

Routine maintenance 31
Periodic checks 31

Cleaning the FRx 32
Disposing of the FRx 32
Ready light troubleshooting tips 32
Troubleshooting a chirping FRx 33

A Accessories 35

B Glossary of terms 37

C Glossary of symbols/controls 41

D Technical information 49

E Configuration 59

F Testing and troubleshooting 65

G Electromagnetic compatibility and immunity information 71

H Shock cycle timing 77

1 INTRODUCTION TO THE HEARTSTART FRx

DESCRIPTION

The Philips HeartStart FRx Defibrillator (861304) is a small, lightweight, portable, rugged, and battery powered automated external defibrillator (AED) indicated to treat victims of sudden cardiac arrest (SCA). The FRx is highly configurable for local protocol considerations.*

The FRx is a public access AED. Users should have received training in basic life support/AED, advanced life support, or a physician-authorized emergency medical response training program.

SUDDEN CARDIAC ARREST

The FRx is used to treat ventricular fibrillation (VF), a common cause of sudden cardiac arrest (SCA), and pulseless ventricular tachycardias (VTs). SCA is a condition that occurs when the heart unexpectedly stops pumping. SCA can occur to anyone – young or old, male or female – anywhere, at any time. Many victims of SCA do not have warning signs or symptoms. Some people may have a higher risk for SCA than others. Causes vary and may be different for infants and children than for adults.

VF is a chaotic quivering of the heart muscle that prevents it from pumping blood. The only effective treatment for VF is defibrillation. The FRx treats VF by sending a shock across the heart, so it can start beating regularly again. Unless this is successful within the first few minutes after the heart stops beating, the victim is not likely to survive.

* Configurability includes timing of the "Call Emergency Medical Services" reminder, CPR protocol variations, and other features. See Appendix E, "Configuration," for details.

INDICATIONS FOR USE

The HeartStart FRx Defibrillator (Model 861304) is indicated for use on potential victims of sudden cardiac arrest (SCA) with the following symptoms:

- Unconsciousness, and
- Absence of normal breathing

The HeartStart FRx (Model 861304) is indicated for adults over 55 pounds (25 kg). The Model 861304 is also indicated for infants and children under 55 pounds (25 kg) or 0-8 years old when used with the optional Infant/Child Key (Model 989803139311). If the Infant/Child Key is not available, or you are uncertain of the child's age or weight, proceed using adult treatment without the infant/child key.

CONTRAINDICATIONS

The HeartStart FRx Defibrillator should not be used when a patient is conscious or breathing normally.

DANGERS, WARNINGS, AND CAUTIONS

It is important to understand how to use your HeartStart FRx Defibrillator safely. Not following or considering this information could lead to a delay of therapy for the patient or cause harm to yourself and others around you. Please read these dangers, warnings, and cautions carefully.

DANGER - Immediate hazards that will result in serious personal injury or death to the user and/or the victim.

WARNING - Conditions, hazards, or unsafe practices that can result in serious personal injury or death.

CAUTION - Conditions, hazards, or unsafe practices that can result in minor personal injury, damage to the FRx Defibrillator, or loss of data stored in the device.

NOTE: The HeartStart FRx Defibrillator is designed to be used only with Philips-approved accessories. The FRx may perform improperly if non-approved accessories are used.

DANGERS

flammable gases If the FRx is used to give a shock in the presence of flammable gases such as in an oxygen tent, there is a risk of explosion. Move supplemental oxygen and oxygen delivery devices away from the defibrillation pads. (However, it is safe to use the FRx on someone wearing an oxygen mask.)

battery The HeartStart M5070A and 989803139301 batteries are not rechargeable. Do not try to recharge, open, crush, or burn the battery, or it may explode or catch fire.

WARNINGS

fluids Do not let fluids get into the FRx. Avoid spilling any fluids on the FRx or its accessories. Spilling fluids into the FRx may damage it or cause a fire or shock hazard. Do not sterilize the FRx or its accessories.

accessories Using damaged or expired equipment or accessories may cause the HeartStart FRx Defibrillator to perform improperly, and/or injure the victim or the user.

patient handling Performing CPR or otherwise handling or moving the patient while the FRx is analyzing heart rhythm can cause an incorrect or delayed analysis. If the FRx tells you a shock is advised while you are handling or moving the patient, stop the vehicle or CPR and keep the patient as still as possible for at least 15 seconds. This will give the FRx time to reconfirm the analysis before telling you to press the Shock button.

The FRx delivers up to 150 joules of electrical energy. This electrical energy may induce ventricular fibrillation or a non-perfusing rhythm in the user or a bystander if the FRx is not used as described in this manual. Improper use of this device can cause serious injury or death. Ensure that the user and any bystanders are not touching the patient when the Shock button is pressed.

children Applying defibrillation to children when it is not necessary may unnecessarily injure a child. Most cardiac arrests in children are not caused by heart problems.

When responding to cardiac arrest in an infant or child:

- Provide infant/child CPR while a bystander calls EMS and retrieves the FRx.

- If no bystander is available, provide 1-2 minutes of CPR before calling EMS and retrieving the FRx.
- If you witnessed the child's collapse, call EMS immediately and then get the FRx.

Alternatively, follow your local protocol.

Keep the FRx out of reach of children to avoid the potential risk of inhalation or swallowing of small parts or strangulation by pads cables.

proximity to other equipment, mobile phones and radios

Use of the FRx adjacent to or stacked with other equipment should be avoided because it could result in improper operation. If such use is necessary, the FRx and the other equipment should be observed to verify that they are operating normally.

The FRx can work correctly when it is fairly close to RF portable communication equipment such as emergency two-way radios and mobile phones, but the communication equipment including peripherals such as antenna cables and external antennas should be used no closer than 12 inches (30 cm) to any part of the FRx. Otherwise, degradation of the performance of the FRx could result. Normally, using a mobile phone near the patient should not cause a problem for the FRx. However, it is best to keep such equipment only as close as necessary to the patient and the FRx.

battery

Removing the battery may reset the FRx, delaying patient defibrillation. Removing and reinserting the battery one or more times when the FRx emits a series of triple chirps may reset the device and cause it to report it is ready for use, though it may be unable to deliver therapy during a rescue. Removing and reinserting the battery when your FRx is emitting a pattern of triple chirps should only be done during an emergency. If your device is emitting a series of triple chirps in stand-by mode, or after an emergency, please remove the FRx from service and contact Philips immediately.

shock hazard

The FRx is not protected from electrical shock hazard if it is opened. The FRx is protected from electrical shock hazard while it is intact. Do not open the FRx, remove its covers, or attempt to repair it. There are no user-serviceable components in the FRx. If repair is required, return the FRx to an authorized service center.

CAUTIONS

device mishandling	The FRx was designed to be sturdy and reliable for many different use conditions. However, handling the FRx too roughly can damage it or its accessories and will invalidate the warranty. Check the FRx and accessories regularly for damage, according to directions.
infectious substances	Avoid directly handling the FRx or its accessories if they have been in contact with body fluids from the patient. Such substances could contain infectious material causing the spread of harmful diseases. Refer to this manual's section on cleaning the FRx after a use if necessary.
maintenance	Improper maintenance may damage the FRx or cause it to function improperly. Maintain the FRx according to directions. Check supplies, accessories, packaging, and spares for damage and expiration dating.
skin burns	Do not let the pads touch each other or other electrodes, lead wires, dressings, medicine patches, etc. In addition, make sure the pads are not in contact with metal objects such as a bed frame or stretcher. Such contact can cause electrical arcing and skin burns during a shock and may also divert the electrical current away from the patient's heart. During a shock, air pockets between the skin and pads can cause skin burns. To help prevent air pockets, make sure pads stick well to the skin. Do not use dried out pads because they will not provide good contact with the skin.
patient handling	There is an electrical shock hazard, or equipment damage hazard if the patient remains connected to devices other than the FRx. Before delivering a shock, it is important to disconnect the patient from other medical electrical equipment, such as blood-flow meters, that may not incorporate defibrillation protection.
patient	The device may be unable to deliver effective defibrillation shocks if the defibrillation pads are not making good contact with the patient's skin. If the patient's chest is not clean, the defibrillation pads may not make adequate contact with the patient. Remove any medicine patches and residual adhesive from the patient's chest before applying the pads.
radiated emissions	While the FRx complies with radiated emission standards, some medical equipment may still be impacted by emissions from the FRx. If this occurs, move the impacted equipment away from the FRx until the FRx is no longer needed for the patient, or EMS arrives and takes over the scene.
environmental conditions	Environmental conditions may cause improper operation. Using the FRx outside of the specified environmental range (temperature, humidity, atmospheric pressure) may result in incorrect or intermittent operation. Make sure the FRx is stored in an environment according to the owner's manual.

configuration	An incorrectly configured language may prevent the FRx from being applied properly. The FRx uses lighted buttons and voice instructions to guide a user through a rescue. If a user is not familiar with the language that the FRx is set to, the FRx may not be used effectively to treat a patient in need, reducing the likelihood of survival for that patient. Make sure that the language of the FRx is set to a language that the majority of the users of that FRx may be familiar with.
pacemakers	The FRx may be unable to deliver effective defibrillation shocks because of an implanted pacemaker in the patient. Do not place the pads directly over an implanted pacemaker or defibrillator. A noticeable lump with a surgical scar should indicate the position of an implanted device.
pads	If the pads do not stick well to the skin, check that the pads adhesive has not dried out. Each pad has a layer of adhesive gel. If the gel is not sticky to the touch, replace the pads with a new set. (For ease of handling, the pad is designed with a non-gel area around the connector cable.)
device operation	Defibrillation may not be provided effectively if an operator delays pressing the Shock button. The FRx will only deliver a shock if the flashing orange Shock button is pressed when the instruction is given. If the Shock button is not pressed within 30 seconds after the instruction, the FRx will disarm itself, and (for the first CPR interval) give a reminder to make sure emergency medical services have been called. The FRx will then begin a CPR interval. This is designed to minimize interruption of CPR and help ensure ongoing patient support.
accessories	The FRx may not be ready for use when needed on a patient if the accessories are not stored correctly. Do not leave the FRx without a set of pads connected; the FRx will start chirping and the i-button will start flashing. The FRx runs daily self-tests. As long as the green Ready light is blinking, it is NOT necessary to test the defibrillator by initiating a battery insertion self-test. This uses battery power and risks draining the battery prematurely.
cpr	Even when CPR is applied correctly, the patient's chest may become bruised or ribs may be fractured. Performing CPR incorrectly could cause additional injuries to a patient, or may not provide needed benefit to the patient. Be sure to follow the CPR guidance instructions provided by the FRx. Be sure to push the i-button to turn on CPR guidance if you are unsure of your CPR skills.
patient handling	Keep the patient still and keep any movement around the patient to a minimum during rhythm analysis. Do not touch the patient or the pads while the Caution light is on solid or flashing. If the FRx is unable to analyze due to electrical "noise" (artifact), it will tell you to stop all movement and remind you not to touch the patient. If the artifact continues for more than 30 seconds, the FRx

will pause briefly to allow you to deal with the source of the noise, then resume analysis.

PROBABLE ADVERSE EFFECTS OF THE DEVICE ON HEALTH

Following are the probable adverse effects (e.g., complications) of the device on health:

- Failure to identify shockable arrhythmia
- Failure to deliver a defibrillation shock in the presence of ventricular fibrillation (VF) or pulseless ventricular tachycardia (VT) which may result in death or permanent injury
- Inappropriate energy, which could cause failed defibrillation or post-shock dysfunction
- Myocardial damage
- Fire hazard in the presence of high oxygen concentration or flammable anesthetic agents
- Incorrectly shocking a pulse sustaining rhythm and inducing VF or cardiac arrest
- Bystander shock from patient contact during defibrillation shock
- Interaction with pacemakers
- Skin burns around the defibrillation pads placement area
- Allergic dermatitis due to sensitivity to the materials used in the defibrillation pads construction
- Minor skin rash

CLINICAL SUMMARY OF SAFETY & PERFORMANCE DATA

Philips, or its predecessor Heartstream, was directly responsible for the conduct of clinical trials related to the safety and effectiveness of the Philips family of AEDs.

ADULT DEFIBRILLATION WAVEFORM

The pivotal clinical trial supporting the Philips SMART biphasic waveform was comprised of three studies. The first was a single center feasibility trial

(Gemini I); followed by a prospective randomized clinical trial (Gemini II), and finally a safety sub-study (Gemini Safety). These studies supported the safety and effectiveness of the SMART biphasic defibrillation waveform.

1. **Gemini I Feasibility Study**[*]

Objective - Gemini I was a clinical evaluation of the transthoracic defibrillation effectiveness of two different biphasic truncated exponential waveforms (115 J and 130 J) with that of a then-standard 200 joules monophasic damped sine waveform.

Study Design - The study was a single site, prospective, randomized and blinded study involving patients undergoing transvenous Implantable Cardioverter Defibrillator (ICD) surgery. Transthoracic ventricular defibrillation rescue shocks were tested after a failed transvenous defibrillation shock was delivered in the course of ICD testing. Each of the three different rescue shocks was tested in random order in each patient. The shock was considered a success if it defibrillated a patient. The biphasic waveforms were generated using a custom, experimental defibrillation (Heartstream) system. The damped sine wave was from the Physio-Control LIFEPAK 6s defibrillator.

Results - Thirty-three patients were enrolled and 30 completed the protocol. Of the 30 patients, 22 were men. All were undergoing a planned procedure for ICD implantation and consented to inclusion in the clinical study. All three waveforms were equally effective at 97%, with one patient failing to be defibrillated with each waveform. The defibrillation energy for the two biphasic waveforms was significantly lower as compared to the damped sine wave ($p < 0.001$), as was the peak current and voltage.

Conclusion - The results showed that biphasic truncated transthoracic shocks of low energy (115 J and 130 J) were as effective in the tested group as 200 joules damped sine wave shocks used in standard transthoracic defibrillators.

* Bardy GH, Gliner BE, Kudenchuk PJ, Poole JE, Dolack GL, Jones GK, Anderson J, Troutman C, Johnson G: Truncated biphasic pulses for transthoracic defibrillation. *Circulation* 1995, 91(6):1768-1774

2. Gemini II Pivotal Study*

Objective - The objective of this randomized, controlled, multi-center trial was to evaluate the safety and effectiveness of the investigational biphasic truncated exponential waveform vs. the control monophasic damped sinusoidal waveform from standard commercially marketed external defibrillators.

Study Design - The study was a prospective, randomized, double-blinded investigation conducted at 14 sites in the United States and Canada. The study population consisted of 318 patients undergoing testing for insertion of an implantable defibrillator or follow-up electrophysiological evaluation post-implantation. In this study rescue shocks of investigational biphasic waveforms of 115 J and 130 J were compared to monophasic waveforms of 200 J and 360 J.

Results - A total of 318 patients were enrolled in the study, and after exclusion criteria were applied there were 294 patients included in the study analyses, for a total of 513 shocks delivered during the study.

Overall, for the 294 included patients analyzed, 513 transthoracic defibrillation attempts (shocks) were performed. The overall breakdown by waveform and success rates is as follows in the table below.

Successful Defibrillations by Waveform Type

Waveform	Successful Defibrillation N(%)	95% Confidence Interval (%)
115 J Biphasic	86 (89)	82-95
130 J Biphasic	144 (86)	81-92
200 J Damped Sine	143 (86)	81-91
360 J Damped Sine	80 (96)	92-100

Conclusion - For the primary hypothesis, the effectiveness of the 130 J truncated biphasic waveform and the 200 J monophasic waveform was not

* Bardy GH, Marchlinski FE, Sharma AD, Worley SJ, Luceri RM, Yee R, Halperin BD, Fellows CL, Ahern TS, Chilson DA et al: Multicenter comparison of truncated biphasic shocks and standard damped sine wave monophasic shocks for transthoracic ventricular defibrillation. Transthoracic Investigators. *Circulation* 1996, 94(10):2507-2514.

significantly different using the Pearson chi-square test (p = 0.97). The 115 J and 130 J biphasic waveforms both demonstrate transthoracic defibrillation effectiveness equivalent to either the 200 J or 360 J monophasic waveforms.

The energy dose increased to 150 J in later clinical studies (ORCA study by Schneider), and 150 J is the energy dose in the SMART biphasic waveform used in the FRx AED.

3. Gemini II Safety Sub-Study*

A single center, prospective analysis was conducted to look at potential differences in ECG ST-segment changes when comparing the waveforms from the pivotal trial. In this study the ST-segment changes were used as a surrogate for myocardial injury. Each patient received two low-energy biphasic waveform shocks at 115 J and 130 J and a 200 J monophasic shock. ECGs were reviewed by two blinded, independent reviewers. This 30-patient sub-study showed that ST-segment elevation was significantly greater for the 200 J damped sine wave ($p<0.001$), indicating a potential safety advantage associated with the biphasic waveform.

4. ORCA (Out of Hospital Response to Cardiac Arrest) Trial†

This postmarket study supports the safe and effective use of the Philips FRx in out-of-hospital defibrillation. The ForeRunner device used in this study, and the FRx device subject to Premarket Approval (PMA), both use SMART biphasic waveforms and PAS shock advisory algorithm technology.

Study Design - Patients were prospectively enrolled in four European EMS systems and included a total of 338 patients. First responders used either impedance-compensating biphasic waveform AEDs (Philips ForeRunner 150 J) or standard monophasic damped sine (MDS) and monophasic truncated exponential (MTE) AEDs with an escalating energy protocol on victims of

* Reddy RK, Gleva MJ, Gliner BE, Dolack GL, Kudenchuk PJ, Poole JE, Bardy GH: Biphasic transthoracic defibrillation causes fewer ECG ST-segment changes after shock. Annals of emergency medicine 1997, 30(2):127-134

† Schneider T, Martens PR, Paschen H, Kuisma M, Wolcke B, Gliner BE, Russell JK, Weaver WD, Bossaert L, Chamberlain D: Multicenter, randomized, controlled trial of 150 J biphasic shocks compared with 200-360 J monophasic shocks in the resuscitation of out-of-hospital cardiac arrest victims. Optimized Response to Cardiac Arrest (ORCA) Investigators. Circulation 2000, 102(15):1780-1787

sudden collapse when defibrillator application was indicated. A sequence of up to three defibrillation shocks was delivered (150 J for each of the three biphasic shocks; for monophasic AEDs, 200 J, 200 J, then 360 J).

Results - A total of 338 patients were enrolled. After exclusion criteria were applied, 115 patients were included in the principal analyses, 54 treated with biphasic and 61 with monophasic AED shocks.

53 of 54 (98%) VF patients were defibrillated using 150 J biphasic shocks compared with 42 of 61 (69%) with 200-360 J monophasic shocks ($p < 0.0001$). The impedance-corrected biphasic truncated exponential (ICBTE) waveform was more effective than the MDS waveform (98% vs. 77%, $p = 0.02$).

A higher percentage of patients (76%) achieved Return of Spontaneous Circulation (ROSC) following 150 J biphasic waveform defibrillation compared with higher energy monophasic waveform defibrillation (54%) ($p = 0.01$).

Conclusion - The study demonstrated that an appropriately dosed low-energy impedance-compensating biphasic waveform strategy results in superior defibrillation performance when compared with escalating, high-energy monophasic shocks in out-of-hospital cardiac arrest. Although survival rates to hospital admission and discharge did not differ, discharged patients who had been resuscitated with biphasic shocks were more likely to have good cerebral performance.

PEDIATRIC DEFIBRILLATION WAVEFORM

Pediatric defibrillation effectiveness is supported with an animal study for the biphasic waveform energy of 50 J and a post-market surveillance study for Pediatric AED use.

1. **Animal Study**[*]

Tang et al. conducted an evaluation of a 50 J biphasic waveform in a porcine model to determine if 50 J was an appropriate energy level (Phase 1), and then

* Tang W, Weil MH, Jorgenson D, Klouche K, Morgan C, Yu T, Sun S, Snyder D: Fixed-energy biphasic waveform defibrillation in a pediatric model of cardiac arrest and resuscitation. Critical care medicine 2002, 30(12):2736-2741

to evaluate the implementation of reducing the adult dose to the pediatric dose by means of attenuated pediatric pads (Phase 2).

In Phase 1 of the Tang et al. study, 4 groups of 5 anesthetized mechanically ventilated piglets weighing 3.8, 7.5, 15, and 25 kg were evaluated for a total of 20 animals. Ventricular fibrillation was induced and after 7 minutes of untreated VF, defibrillations were attempted with an impedance-compensating biphasic waveform defibrillator modified to deliver shocks with a nominal energy level of 50 J.

In Phase 2 of the study, shocks were delivered through special pediatric pads in conjunction with a conventional adult AED (FR2; Philips Medical Systems). The same VF induction and resuscitation protocol as Phase 1 was exercised on 3 piglets in 3 weight groups (3.7, 13.5, and 24.2 kg). The SMART biphasic waveform as implemented in the FR2 used in this study supports the SMART biphasic waveform as implemented on the FRx AED.

In both phases, all animals were successfully resuscitated. The average total number of shocks and total delivered energy was not weight dependent ($p < 0.05$). Post-resuscitation hemodynamic and myocardial function quickly returned to baseline values in both experimental groups; 100% of the animals survived. Animals were monitored for survival at 24, 48 and 72 hours in Phase 1 and in Phase 2 for four hours; all animals survived through the last time-point.

2. Postmarket Surveillance Study of Pediatric AED Use*

The objective of the postmarket surveillance study was to confirm that certain adult AEDs with shock intensity attenuation could be used safely and effectively in the pediatric population. The study population was infants and children less than 8 years of age or under 55 pounds. The HeartStart FR2 Defibrillator is a predecessor to the FRx AED. Data from this defibrillator is applicable to the safety and effectiveness of the FRx AED.

Study Design - This prospective, observational, postmarket surveillance study included the Philips FR2 AED and Pediatric Attenuated Electrodes, and the HeartStart OnSite AED with attenuation pads cartridge. Data from the FR2 and

* Atkins DL, Jorgenson DB: Attenuated pediatric electrode pads for automated external defibrillator use in children. Resuscitation 2005, 66(1):31-37

HeartStart OnSite are applicable to the consideration of the safety and effectiveness of the FRx because the FRx AED uses the same principles for its SMART biphasic therapy waveform and PAS patient analysis algorithm.

Results - Through September 2004, there were 26 confirmed pediatric-use cases: 25 of 26 reported use of the FR2 and one reported use of the OnSite. The median age was 2 years. The users were predominately EMS personnel or health care professionals (n=24). Most arrests occurred at home (n=16). Most patients to whom the device was applied had non-shockable rhythms (16, of which 13 were confirmed with AED data). Of 7 patients who had ventricular fibrillation and received attenuated shocks, all had termination of ventricular fibrillation and 5 survived to hospital discharge. The median age of the 7 patients was 3 years (range 18 months to 10 years). These patients received on average 2 shocks (range 1-4).

Conclusion - Based on the postmarket surveillance data, the FR2 AED used with the FR2 infant/child attenuated pads and the HeartStart OnSite AED used with infant/child pads cartridge performed safely and effectively in the pediatric population, which can be applied to the pediatric use of the FRx.

ESSENTIAL PERFORMANCE

The HeartStart FRx Defibrillator maintains safe and effective performance of the defibrillation therapy functions when operated in the electromagnetic environment specified in the Appendix G tables.

PRINCIPLES OF OPERATION

The HeartStart FRx Defibrillator is designed to provide external defibrillation therapy to someone experiencing sudden cardiac arrest caused by ventricular fibrillation (VF), the most common cause of SCA, and pulseless ventricular tachycardias (VTs). The only effective treatment for these non-perfusing arrhythmias is defibrillation. The FRx treats them by sending a shock across the heart, so it can start beating regularly again. The FRx is designed to be easy to use. In its default AED mode, when connected to defibrillator pads that are properly applied to the patient's bare chest, the FRx prompts you to take specific actions; automatically analyzes the patient's heart rhythm and advises you whether or not the rhythm is shockable; and, if advised by its rhythm

analysis algorithm, arms the Shock button and instructs you to press it to deliver a biphasic electric pulse designed to defibrillate the heart. For detailed instructions for use, see Chapter 3, "Using the HeartStart FRx."

IMPLEMENTATION CONSIDERATIONS

Check with your local health department to see if there are any national or local requirements about owning and using a defibrillator. The HeartStart FRx Defibrillator is one part of a well-designed emergency response plan. Recognized resuscitation councils recommend that emergency response plans include physician oversight and training in cardiopulmonary resuscitation (CPR).

Several national and local organizations offer combined CPR/AED training. Philips recommends that you train on the device you will be using. Contact your Philips representative for information, or visit us online at www.philips.com/AEDsupport.

NOTE: Training accessories are available for practicing use of the FRx. See Appendix A for information.

FOR MORE INFORMATION

Contact your local Philips distributor for additional information about the FRx. We will be happy to answer any questions you may have and to provide you with copies of the clinical summaries of several key studies using Philips automated external defibrillators.*

* Clinical summaries also include defibrillators sold as ForeRunner and FR2.

2 SETTING UP THE HEARTSTART FRx

PACKAGE CONTENTS

Check the contents of the FRx box to be sure it contains:

- 1 HeartStart FRx Defibrillator
- 1 four-year battery M5070A,* pre-installed
- 1 package of HeartStart SMART Pads II 989803139261, containing one set of adhesive defibrillation pads in a disposable plastic case, pre-installed
- 1 Quick Reference Guide
- 1 Owner's Manual
- 1 HeartStart Quick Setup Guide
- 1 inspection log/maintenance booklet with plastic storage sleeve and maintenance tags

IMPORTANT NOTE: The FRx is designed to be used with a carry case. A number of carry cases are offered to meet the needs of your individual defibrillation program. These include a standard carry case and a hard-shell carry case. See Appendix A for information about these as well as a list of training materials and other accessories available from Philips.

If you have purchased the FRx Ready-Pack configuration, the FRx is installed in an FRx carry case, which also contains a spare SMART Pads II case.

SETTING UP THE FRx

Setting up the FRx is simple and quick. The Quick Setup Guide provides illustrated instructions for setting up the FRx, described in detail below.

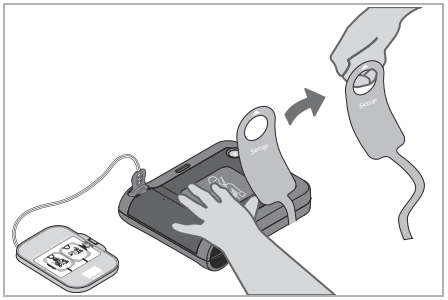
1. Remove the FRx from its packaging. Check that the battery and SMART Pads II are installed.†

* The FRx sold for aviation applications includes a TSO-certified battery, REF: 989803139301, instead of the M5070A.

† If the battery and pads are not installed, follow the directions in Chapter 4, "After using the HeartStart FRx", to install the pads and battery.

NOTE: To prevent the pads' adhesive gel from drying out, do not open the SMART Pads II case until you need to use the pads.

2. Pull out and discard the green Setup tab.



3. The FRx will automatically run a self-test. Press the Shock button and the On/Off button when instructed. Be sure to let the self-test run all the way to completion. When the self-test is over, the FRx will report the result, and tell you to push the green On/Off button in case of an emergency. (*Do not push the green button unless this is an actual emergency.*) Then the FRx will turn off and go to standby mode.* The green Ready light will be blinking to show the FRx is ready for use.
4. Install the FRx in its carry case, if it is not pre-installed. Check that the Quick Reference Guide† is face up in the clear plastic window on the inside of the carry case. Philips recommends that you store a spare pads case and spare battery with your FRx. If you are using an FRx carry case, there is an area in the lid of the case, under the flap, to store a spare package of pads and a spare battery.‡

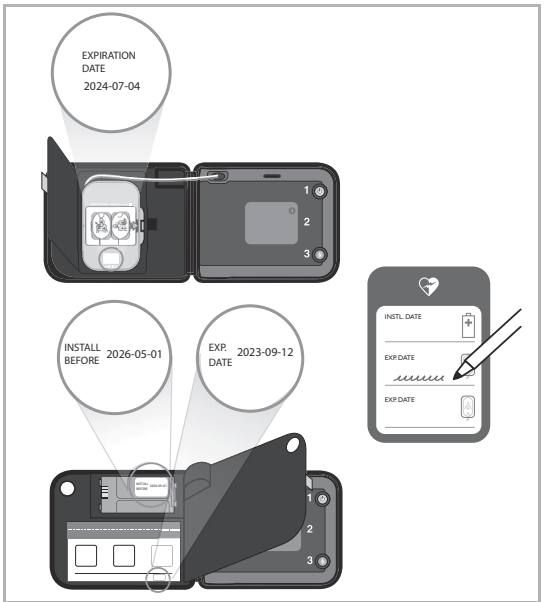
NOTE: Do not store anything in the defibrillator carry case that it is not designed to accommodate. Store all objects in their intended location in the case.

* As long as a battery is installed, turning the FRx "Off" puts it into standby mode, which means that it is ready for use.

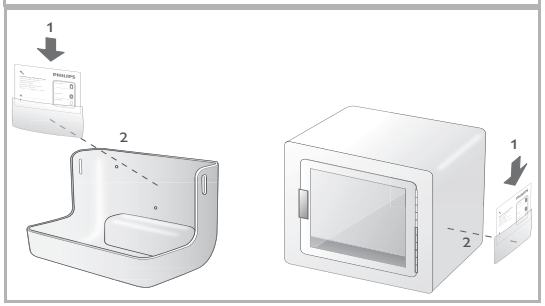
† The illustration on the cover of the Quick Reference Guide is a 3-step guide to using the FRx. Detailed illustrated directions are inside, for reference in an emergency, or if you are hearing impaired or using the FRx where it is hard to hear the voice instructions. Any of the carry case options has room for storing the Quick Reference Guide.

‡ See Chapter 4, "After using the HeartStart FRx" for directions on how to replace the battery in the FRx.

5. Use the maintenance tag provided to record the expiration date of the installed pads. If you have a spare pads case and spare battery, record the pads expiration date and battery install-by date on the maintenance tag.



6. The maintenance tag and maintenance booklet should be kept with your FRx. Adhere the plastic storage sleeve for the booklet to the AED wall mount or cabinet and place the booklet in it.



7. Store the FRx in accordance with your site's emergency response protocol. Typically, this will be in a high-traffic area that is easy to access, convenient for checking the Ready light periodically, and easy to hear the alarm chirp if the battery power gets low or the defibrillator needs attention. Ideally, the FRx should be stored near a telephone, so the Emergency Response Team or Emergency Medical Services can be alerted as fast as possible in the event of a possible SCA. Ensure storage area is free of pests and not accessible to pets or children. The FRx is also transportable in a road ambulance or fixed-wing aircraft.

In general, treat the FRx as you would any piece of electronic equipment, such as a computer. Be sure to store the FRx according to its specifications. See Appendix D for details. As long as a battery and a pads set are installed, the green Ready light should be blinking to show that the FRx has passed its most recent self-test and is therefore ready to use.

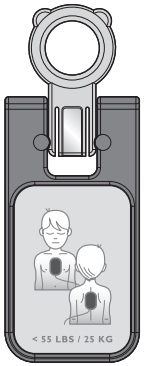
NOTE: Always store the FRx with a set of SMART Pads II and a battery installed, so it will be ready to use and can perform daily self-tests. Training Pads II should be stored separately from the FRx to avoid confusion during use.

RECOMMENDED ACCESSORIES

It is always a good idea to have a spare battery and a spare pads set. Other things that are useful to keep with the FRx include:

- scissors — for cutting the victim's clothes if needed
- disposable gloves — to protect the user
- a disposable razor — to shave the chest if hair prevents good pads contact
- a pocket mask or face shield — to protect the user
- a towel or absorbent wipes — to dry the victim's skin for good pads contact

Philips has a Fast Response Kit with all these items. See Appendix A for information.



Infant/Child Key accessory.

If you may need to defibrillate an infant or a child under 55 pounds (25 kg) or 0-8 years old, it is recommended that you order the Infant/Child Key accessory, available separately. When the Infant/Child Key is inserted in the FRx, the FRx automatically reduces the defibrillation energy to 50 joules and, if optional CPR guidance is selected, provides guidance appropriate for infants and children. Directions for using the Infant/Child Key are provided in Chapter 3, "Using the HeartStart FRx".

See Appendix A for a list of accessories and training products for the FRx available from Philips.

3 USING THE HEARTSTART FRx

IMPORTANT NOTE: Be sure to read the DANGERS, WARNINGS, and CAUTIONS throughout the Owner's Manual as well as in Chapter 1.

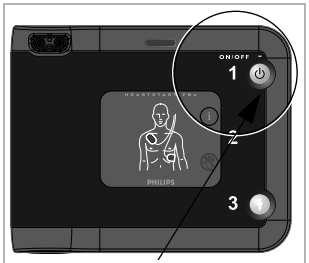
OVERVIEW

If you think someone is in SCA, act quickly and calmly.

- If the patient is an infant or child, first perform CPR, then call for emergency medical services (EMS) before you apply the FRx. See the special section on treating infants and children.
- Call your emergency services provider. If someone else is available, ask him or her to call for emergency medical assistance while you get the FRx.
- Quickly get the FRx and bring it to the victim's side. If there is any delay in getting the defibrillator, check the patient and perform cardiopulmonary resuscitation (CPR) if needed until the FRx is available.
- It is safe to use the HeartStart FRx Defibrillator on a patient lying on a wet surface. Before doing so, remove the patient from standing water, such as a pool or bathtub. It is also safe to use the HeartStart FRx Defibrillator on a patient lying on a conductive surface, such as a metal surface. It is important to dry the patient's chest completely, so that the pads stick well to the dry, bare skin.
- Check the immediate environment for flammable gases. Do not use the FRx in the presence of flammable gases, such as an oxygen tent. However, it is safe to use the FRx on someone wearing an oxygen mask.

There are three basic steps to using the defibrillator to treat someone who may be in sudden cardiac arrest:

1. Press the green On/Off button.
2. Follow the FRx voice instructions.
3. Press the flashing orange Shock button if instructed.

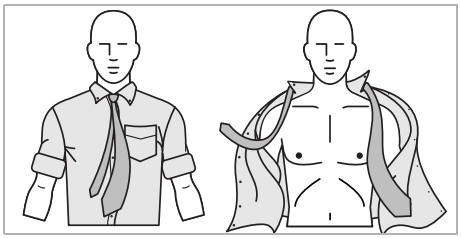


Press green On/Off button.

STEP 1: PRESS THE GREEN ON/OFF BUTTON

Press the On/Off button ⏻ to turn on the FRx.

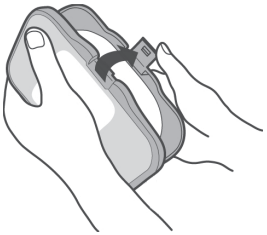
The FRx tells you to remove all clothes from the person's chest. If necessary, rip or cut off the clothing to bare the person's chest.

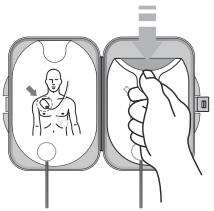


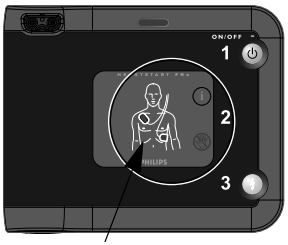
STEP 2: FOLLOW THE FRx VOICE INSTRUCTIONS

Remove the SMART Pads II case from the carry case. Clean and dry the patient's skin, and, if necessary, clip or shave excessive chest hair to ensure good pads contact with the bare skin.

Open the pads case as shown below. Peel off one pad.


Open pads case.

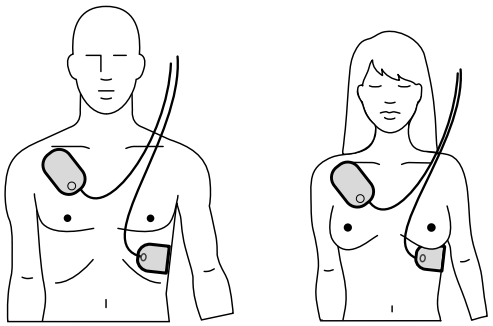

Peel one pad from case.



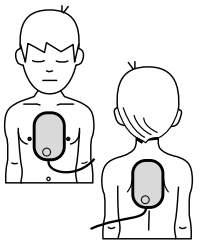
Pads icons flash.

Pads placement is very important. The icons on the pads placement diagram on the FRx front panel will be flashing, to help guide you. Place the pad on the patient's bare skin *exactly as shown* in the following drawing. Press the adhesive portion of the pad down firmly. Then repeat this with the other pad.

Where to place pads on adults (anterior-anterior).



Where to place pads on infants or children under 55 pounds (25 kg) or 0-8 years old (anterior-posterior).



STEP 3: PRESS THE FLASHING ORANGE SHOCK BUTTON IF INSTRUCTED

As soon as the FRx detects that the pads are attached to the patient, the pads icons turn off. The FRx begins analyzing the patient's heart rhythm. It tells you that no one should be touching the patient, and the Caution light begins flashing as a reminder.



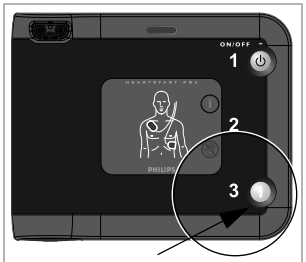
Caution light

If a shock is needed:

The Caution light stops flashing and stays on, and the orange Shock button starts flashing. The FRx tells you to press the flashing orange button. You must press the Shock button for a shock to be delivered. Before you press the button, make sure no one is touching the patient. When you press the Shock button, the FRx tells you that the shock has been delivered. Then the defibrillator tells you it is safe to touch the patient, instructs you to begin CPR, and invites you to press the flashing blue i-button for CPR guidance if desired.



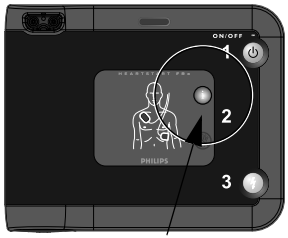
Shock button



Shock button flashes.

If a shock is not needed:

The i-button comes on solid, to show that it is safe to touch the patient. The FRx also tells you to perform CPR if needed. (If CPR is not needed – for example, if the patient is moving or regaining consciousness – follow your local protocol until emergency medical personnel arrive.) Then the FRx invites you to press the flashing blue i-button for CPR guidance if desired.



i-button flashes.

For CPR guidance:

Press the flashing blue i-button during the first 30 seconds of the patient care pause to activate CPR guidance.* (If the Infant/Child Key is inserted, the CPR guidance provided will be for infant/child CPR.) When the pause is over, the defibrillator tells you to stop CPR, so it can analyze the patient's heart rhythm. The motion caused by CPR can interfere with analysis, so be sure to stop all motion when instructed.



i-button

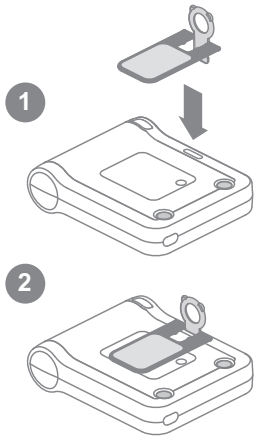
TREATING INFANTS AND CHILDREN

WARNING: Most cardiac arrests in children are not caused by heart problems. When responding to cardiac arrest in an infant or child:

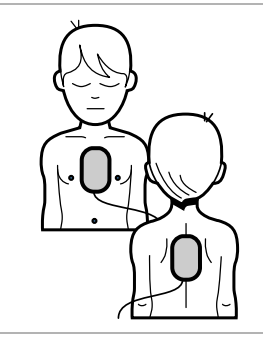
- Provide infant/child CPR while a bystander calls EMS and brings the FRx.
- If no bystander is available, provide 1-2 minutes of CPR before calling EMS and retrieving the FRx.
- If you witnessed the child's collapse, call EMS *immediately* and *then* get the FRx.

Alternatively, follow your local protocol.

* The default configuration for the FRx provides CPR guidance when you press the i-button in this situation; however, the default setting can be revised by your Medical Director using Philips software available separately. See Appendix E for more information.



Insert the Infant/Child Key if available.



If the victim is under 55 pounds (25 kg) or 0-8 years old, and you have an Infant/Child Key:

- Insert the Infant/Child Key into the slot at the top center of the front panel of the FRx (see illustration at left). The pink portion of the Key pivots (1) and fits into the slot (2), with the front of the Key lying flat on the surface of the FRx so the infant/child pads placement diagram is visible. (The back of the Infant/Child Key also has a diagram showing how to insert it.)
- Turn on the FRx and follow instructions to remove all clothing from the torso, to bare both the chest and the back.
- Place the pads on the child's front and back, as illustrated. It does not matter which pad is placed on the chest or the back.

NOTE: It does not matter whether you insert the Infant/Child Key before or immediately after turning on the FRx. However, the Infant/Child Key should be inserted before placing the pads on the patient.

With the Infant/Child Key inserted, the FRx will announce "Infant/Child Mode," automatically reduce the defibrillation energy from the adult dose of 150 Joules to 50 joules,* and provide optional infant/child CPR guidance.

If the Infant/Child Key is removed during use, the FRx will announce "Adult Mode." Any shocks delivered will be at adult energy, and optional CPR guidance will be for adult CPR.

If the victim is under 55 pounds (25 kg) or 0-8 years old, but you do NOT have an Infant/Child Key:

- DO NOT DELAY TREATMENT.
- Turn on the FRx and follow instructions to remove all clothing from the torso, to bare both the chest and the back.
- Place the one pad in the center of the chest between the nipples, and the other in the center of the back (anterior-posterior).

* This lower energy level may not be effective for treating an adult.

If the victim is over 55 pounds (25 kg) or greater than 8 years old, or if you are not sure of the exact weight or age:

- DO NOT DELAY TREATMENT.
- Turn on the FRx without inserting the Key and follow instructions to remove all clothing from the chest.
- Place the pads as illustrated on each pad (anterior-anterior). Make sure the pads do not overlap or touch each other.

WHEN EMERGENCY MEDICAL SERVICES ARRIVE

When Emergency Medical Services (EMS) personnel arrive to care for the patient, they may decide to apply another defibrillator to allow monitoring of the patient. Depending on their equipment, the EMS team may apply different pads. In that case, the SMART Pads II should be removed. EMS personnel may want a summary of the last-use data* stored in the FRx. To hear the summary data, hold down the i-button until the FRx beeps.

NOTE: After the EMS team removes the SMART Pads II from the patient, remove the Infant/Child Key, if used, and install a new SMART Pads II set before returning the FRx to service, to be sure it is ready for use.

REMINDERS

- Remove any medicine patches and residual adhesive from the patient's chest before applying the pads.
- Do not place the pads directly over an implanted pacemaker or defibrillator. A noticeable lump with a surgical scar should indicate the position of an implanted device.
- Do not allow the pads to contact other electrodes or metal parts that are in contact with the patient.
- If the pads do not stick well, check that the pads adhesive has not dried out. Each pad has a layer of adhesive gel. If the gel is not sticky to the touch, replace the pads with a new set. (For ease of handling, the pad is designed with a non-gel area around the connector cable.)

* See Chapter 4, "After Using the HeartStart FRx," for details about data storage.

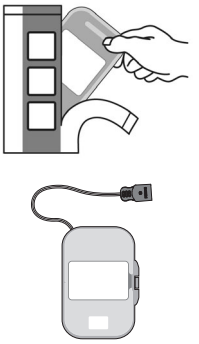
- Keep the patient still and keep any movement around the patient to a minimum during rhythm analysis. Do not touch the patient or the pads while the Caution light is on solid or flashing. If the FRx is unable to analyze due to electrical "noise" (artifact), it will tell you to stop all movement and remind you not to touch the patient. If the artifact continues for more than 30 seconds, the FRx will pause briefly to allow you to deal with the source of the noise, then resume analysis.
- The FRx will only deliver a shock if the flashing orange Shock button is pressed when the instruction is given. If the Shock button is not pressed within 30 seconds after the instruction, the FRx will disarm itself, and (for the first CPR interval) give a reminder to make sure emergency medical services have been called, then begin a CPR interval. This is designed to minimize interruption of CPR and help ensure ongoing patient support.
- While waiting for you to press the Shock button, the FRx will continue to analyze the heart rhythm. If the patient's rhythm changes before you press the Shock button, and a shock is no longer needed, the defibrillator will disarm and tell you a shock is not advised.
- If for any reason you want to turn off the defibrillator during a use, you can press the On/Off button – holding it down for at least one second – to return the device to standby mode.

Intentionally blank.

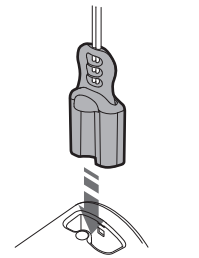
4 AFTER USING THE HEARTSTART FRx

AFTER EACH USE

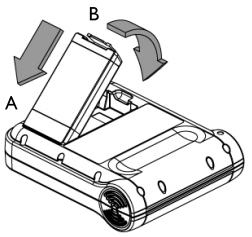
1. Check the outside of the FRx for signs of damage, dirt, or contamination. If you see signs of damage, contact Philips for technical support. If the defibrillator is dirty or contaminated, clean it according to the guidelines in Chapter 5.
2. The single-use SMART Pads II must be replaced after being used. Open the SMART Pads II package and take out the pads case (A). *Do not open the pads case until you need to use the pads in an emergency.*
3. Plug the SMART Pads II cable connector into the connector port on the FRx (B). Store the unopened SMART Pads II case in the pocket provided in the defibrillator carry case.
4. Check supplies and accessories for damage and expiration dates. Replace any used, damaged or expired items. Use a new maintenance tag to record the pads expiration date for the new installed pads. If you replace the spare pads and/or battery be sure to record the dates for them on the maintenance tag as described in Chapter 2. Then sign and date the inspection log/maintenance booklet.
5. Unless your protocol requires that the battery remain installed, remove the battery for five seconds. Then reinstall the battery by placing the bottom end (A) of the battery into the bottom of the compartment on the back of the FRx, then firmly pressing down the top (latch) end of the battery into the compartment, until it clicks into place (B).



A. Remove the pads case from packaging.



B. Plug in the pads cable connector.



Insert battery in the FRx.

6. The FRx will automatically run a self-test when the battery is inserted. Press the Shock button and On/Off button when instructed. Be sure to let the self-test run all the way to completion. When the self-test is over, the FRx will report the result, and tell you to push the green On/Off button in case of an emergency. (*Do not push the green button unless this is an actual emergency.*) Then the FRx will turn off and go to standby mode. The green Ready light will be blinking to show the FRx is ready for use.*

   NOTE: Always store the FRx with a set of SMART Pads II and a battery installed, so it will be ready to use and can perform daily self-tests.

7. Return the FRx to its storage location so it will be ready for use when needed. Place the updated inspection log/maintenance booklet on the defibrillator wall mount or cabinet.

FRx DATA STORAGE

The FRx automatically stores data about its last clinical use in its internal memory. The stored data can be conveniently transferred to a personal computer running the appropriate application in the Philips HeartStart Event Review Pro data management software. Event Review Pro software is for use by trained personnel only. Information about HeartStart Event Review Pro is available online at www.philips.com/eventreview.

* As long as a battery is installed, turning the FRx "Off" puts it into standby mode, which means that it is ready for use.

Follow your local protocol with regard to prompt data transfer for medical review after using the FRx.* Details about data transfer and timing are provided in Event Review Pro documentation.

The information automatically stored by the FRx includes a summary of last-use data and detailed data about its last clinical use. You can get a voice summary of information about the last use of the defibrillator by holding the i-button down until it beeps once. The FRx will tell you how many shocks were delivered and how long it has been since it was turned on. Summary data are available anytime the defibrillator is ready for use (the battery and pads are installed, and the defibrillator is not turned on) or while it is actually in use. Removing the battery erases the summary data for the last use.

Last-use data stored in internal memory include:

- ECG recordings (a maximum of 15 minutes following pads application†)
- the FRx status (entire incident)
- the FRx rhythm analysis decisions (entire incident)
- the elapsed time associated with stored events (entire incident)

* The FRx automatically stores information about its last clinical use in its internal memory for at least 30 days, so the data can be downloaded to a computer running appropriate Event Review Pro software. (If the battery is removed during this period, the defibrillator retains the files. When the battery is reinstalled, the last-use ECG recording will be kept in defibrillator memory for an additional 30 days.) After this time, the last-use ECG recordings will automatically be erased to prepare for a future use.

† If ECG recordings from a previous use have not been erased, the maximum time for new ECG recordings may be less.

Intentionally blank.

5 MAINTAINING THE HEARTSTART FRx

ROUTINE MAINTENANCE

The FRx is very simple to maintain. The defibrillator performs a self-test every day. In addition, a battery insertion self-test is run whenever a battery is installed in the device. The defibrillator's extensive automatic self-test features eliminate the need for any manual calibration.

WARNING: Electrical shock hazard. Do not open the FRx, remove its covers, or attempt repair. There are no user-serviceable components in the FRx. If repair is required, return the FRx to an authorized service center.

REMINDERS:

- Do not leave the defibrillator without a set of pads connected; the defibrillator will start chirping and the i-button will start flashing.
- Do not store the FRx with the Infant/Child Key installed.
- The FRx runs daily self-tests. As long as the green Ready light is blinking, it is NOT necessary to test the defibrillator by initiating a battery insertion self-test. This uses battery power and risks draining the battery prematurely.

PERIODIC CHECKS

Other than the checks recommended after each use of the FRx, maintenance is limited to periodically checking the following:

- Check the green Ready light. If the green Ready light is not blinking, see Troubleshooting Tips, below.
- Replace any used, damaged or expired supplies and accessories.
- Check the outside of the defibrillator. If you see cracks or other signs of damage, contact Philips for technical support.

Record each periodic check in your inspection log/maintenance booklet.

CLEANING THE FRx

The outside of the HeartStart FRx can be cleaned with a soft cloth dampened in soapy water, chlorine bleach (2 tablespoons per quart or liter of water), ammonia-based cleaners, or 70% isopropyl (rubbing) alcohol. It is recommended that the carry case be cleaned with a soft cloth dampened in soapy water.

REMINDERS:

- Do not use strong solvents such as acetone or acetone-based cleaners, abrasive materials, or enzymatic cleaners to clean the FRx and accessories.
- Do not immerse the FRx in fluids.
- Do not sterilize the FRx or its accessories.

DISPOSING OF THE FRx

The FRx and its accessories should be disposed of in accordance with local regulations.

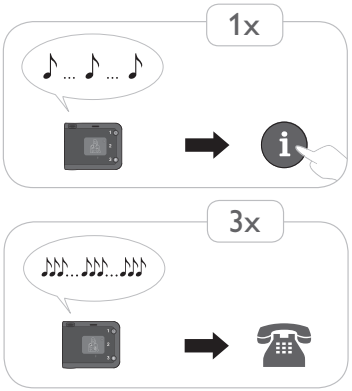
READY LIGHT TROUBLESHOOTING TIPS

The green Ready light is your guide to knowing if the defibrillator is ready for use.

- If the Ready light is blinking: The FRx has passed the battery insertion self-test and the last periodic self-test and is therefore ready for use.
- If the Ready light is solid: The FRx is in use or running a self-test.
- If the Ready light is off, the FRx is emitting a series of single chirps, and the i-button is flashing: A self-test error has occurred, there is a problem with the pads, the Infant/Child Key has been left installed, or the battery power is low. Press the i-button for instructions.
- If the Ready light is off, and the FRx is emitting a series of triple chirps, please call Philips for technical support. See "Troubleshooting a chirping FRx" for more information.
- If the Ready light is off but the FRx is not chirping and the i-button is not flashing: there is no battery inserted, the battery is depleted, or the defibrillator needs repair. Insert/replace battery and run the self-test. As long as the FRx passes the self-test, you can be assured it is ready for use.

TROUBLESHOOTING A CHIRPING FRx

Your FRx tests itself at regular intervals to ensure it is ready for use. If your FRx emits a series of single chirps (♪... ♪... ♪...), press the flashing blue i-button for information.



A triple-chirp alert (♪♪♪... ♪♪♪... ♪♪♪...) could mean that a potentially serious problem was detected during self-test that could prevent your FRx from delivering therapy in an emergency. If you ever hear your FRx emit a series of triple chirps:

- in stand-by mode — please call Philips immediately for technical support at the regional number listed on the back cover of this manual.
- in an emergency rescue — press the flashing blue i-button and follow the voice prompts. Removing and reinserting the battery can clear some errors and equip the device to deliver therapy in a rescue. The battery removal and reinsertion procedure should only be done in an emergency situation. Once the emergency is over, please call Philips immediately for technical support.

WARNING: Removing and reinserting the battery one or more times when an FRx emits a series of triple chirps may reset the device and cause it to report it is ready for use, though it may be unable to deliver therapy during a rescue. Removing and reinserting the battery when your FRx is emitting a pattern of triple chirps should only be done during an emergency. If your device is emitting a series of triple chirps in stand-by mode, or after an emergency, please remove the FRx from service and contact Philips immediately.

More detailed testing and troubleshooting information is available in Appendix F.

A ACCESSORIES

Accessories* for the HeartStart FRx Defibrillator 861304 available separately from your local Philips representative or online at www.philips.com/AED include:

- Batteries (spare is recommended)
  - Battery [REF: M5070A]
  - Aviation applications battery [REF: 989803139301]
- HeartStart SMART Pads II [REF: 989803139261] (spare is recommended)
- Carry Cases
  - FRx carry case [REF: 989803139251]
  - Plastic waterproof hardshell carry case [REF:YC]
- Cabinets and Wall Mounts
  - AED wall mount bracket [REF: 989803170891]
  - Basic surface-mounted cabinet [REF: 989803136531]
  - Premium surface-mounted cabinet [REF: PFE7024D]
  - Premium semi-recessed cabinet [REF: PFE7023D]
- AED Signage
  - AED awareness placard, red [REF: 989803170901]
  - AED awareness placard, green [REF: 989803170911]
  - AED Wall Sign, red [REF: 989803170921]
  - AED Wall Sign, green [REF: 989803170931]
- Infant/Child Key [REF: 989803139311]
- Fast Response Kit (pouch containing a pocket mask, a disposable razor, two pairs of disposable gloves, a pair of paramedic's scissors, and an absorbent wipe) [REF: 68-PCHAT]
- Infrared adapter for use with HeartStart Event Review Pro software [REF: ACT-IR]
- HeartStart FRx Defibrillator Quick Reference Guide [REF: 989803138601]

* Certain accessories require a prescription in the United States.

- Training
  - HeartStart Training Pads II (kit containing one set of Training Pads II in training pads case, adult pads placement guide, Instructions for Use, and illustrated guide) [REF: 989803139271]
  - Replacement Training Pads II (pair of training pads on disposable liner for use in training pads case provided with HeartStart Training Pads II) [REF: 989803139291]
  - Adult pads placement guide [REF: M5090A]
  - Infant/Child pads placement guide [REF: 989803139281]

Data Management Software

- HeartStart Configure [REF: 861487]
- HeartStart Event Review Pro [REF: 861431]
- HeartStart Data Messenger [REF: 861451]

B GLOSSARY OF TERMS

The terms listed in this Glossary are defined in the context of the Philips HeartStart FRx Defibrillator 861304 and its use.

AED	Automated external defibrillator (a semi-automatic defibrillator).
AED mode	The standard treatment mode for the HeartStart FRx Defibrillator. It provides voice instructions guiding the rescuer through applying the adhesive pads, waiting for rhythm analysis, and delivering a shock if needed.
analysis	See "SMART analysis."
arrhythmia	An unhealthy, often irregular, beating of the heart.
artifact	Electrical “noise” caused by sources such as muscle movements, CPR, patient transport, or static electricity that may interfere with rhythm analysis.
battery	The sealed lithium manganese dioxide battery used to power the HeartStart FRx Defibrillator. It is provided in a pack that fits into a compartment on the back of the defibrillator.
caution light	A light on the front of the HeartStart FRx Defibrillator that flashes during rhythm analysis and is on solid when a shock is advised, as a reminder not to touch the patient
configuration	The settings for all operating options of the HeartStart FRx Defibrillator, including treatment protocol. The factory default options can be modified by authorized personnel using HeartStart Configure software.
CPR	Cardiopulmonary resuscitation. A technique for providing artificial respiration and heart compressions.
CPR guidance	Basic verbal instructions for performing cardiopulmonary resuscitation, including hand placement, rescue breathing, compression depth and timing, provided by the FRx when the flashing blue i-button is pressed during the first 30 seconds of a patient care pause.
defibrillation	Termination of cardiac fibrillation by applying electrical energy.
ECG	Electrocardiogram, a record of the electrical rhythm of the heart as detected through defibrillation pads.

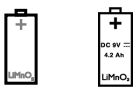
fibrillation	A disturbance of the normal heart rhythm that results in chaotic, disorganized activity that cannot effectively pump blood. Ventricular fibrillation (fibrillation in the lower chambers of the heart) is associated with sudden cardiac arrest.
HeartStart Event Review Pro	A software application used for data management and post event review. Information is available from Philips Medical Systems at www.philips.com/eventreview.
i-button	An "information" button on the front of the HeartStart FRx Defibrillator. If the i-button is pressed during the 30 seconds it flashes during a patient care pause, the FRx provides CPR guidance;* if the i-button is pressed when it is flashing and the FRx is chirping, the FRx provides troubleshooting guidance. At other times, if the i-button is pressed and held until it beeps once, the FRx provides summary information about its last clinical use and device status. When the i-button is on solid (not flashing), it indicates the user may safely touch the patient.
Infant/Child Key	A "key" recommended for use when defibrillating a potential SCA victim under 55 pounds or 0-8 years old. When inserted into a dedicated slot on the FRx's front panel, the Infant/Child Key illustrates correct pads placement, with lighted icons, on these young victims. With the Infant/Child Key inserted, the HeartStart FRx Defibrillator automatically reduces the energy of any shock delivered to 50 joules and provides CPR guidance, if selected, appropriate for infants and children.
infrared (IR) communications	A method of sending information using a special part of the light spectrum. It is used to transmit information between the HeartStart FRx Defibrillator and a computer running HeartStart Event Review Pro software.
NSA	"No Shock Advised," a decision made by the HeartStart FRx Defibrillator that a shock is not needed, based on analysis of the patient's heart rhythm.
NSA pause	A pause provided by the HeartStart FRx Defibrillator following an No Shock Advised (NSA) decision. The pause can be configured to a "standard" NSA pause or a "SMART" NSA pause. During a standard NSA pause the defibrillator performs no background monitoring of the patient rhythm. During a SMART NSA pause, the defibrillator conducts background monitoring and, if it detects an artifact-free shockable rhythm, will exit the pause and begin rhythm analysis. If the HeartStart FRx Defibrillator detects artifact such as that created by CPR, or if the user presses the i-button for CPR guidance during a SMART NSA pause, the defibrillator will not exit the

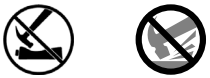
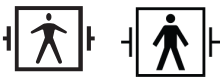
* Pressing the i-button for CPR guidance during a SMART NSA pause turns off background monitoring.

	pause for rhythm analysis in order to allow CPR to be completed uninterrupted.
non-shockable rhythm	A heart rhythm that the HeartStart FRx Defibrillator determines is not appropriate for defibrillation.
On/Off button	A green button located on the front of the HeartStart FRx Defibrillator. Pressing the On/Off button when the defibrillator is in standby mode turns the defibrillator on; pressing and holding the On/Off button for one second when the defibrillator is on turns the defibrillator off and disarms the defibrillator. In addition, pressing the On/Off button stops the battery insertion self-test that automatically runs when a battery is inserted.
pads	See "SMART Pads II."
patient care pause	A defined period to allow CPR. See "NSA pause" and "protocol pause."
periodic self-tests	Daily, weekly, and monthly tests automatically conducted by the HeartStart FRx Defibrillator when it is in its standby mode. The tests monitor many key functions and parameters of the defibrillator, including battery capacity, pads readiness, and the state of its internal circuitry.
protocol	A sequence of operations performed by the HeartStart FRx Defibrillator to direct patient care in the AED mode.
protocol pause	A period provided by the HeartStart FRx Defibrillator after a shock series, during which the responder can administer CPR. The defibrillator does not conduct background monitoring of the patient's heart rhythm during this pause.
Quick Shock	The ability of the FRx to deliver a defibrillation shock very quickly – typically within 8 seconds – after the end of a patient care pause.
Ready light	A green LED showing the readiness for use of the HeartStart FRx Defibrillator. A blinking Ready light means the defibrillator is ready for use; a solid Ready light means the defibrillator is being used.
rhythm analysis	See "SMART analysis."
Shock button	An orange button with a lightning bolt symbol on it, located on the front of the HeartStart FRx Defibrillator. The Shock button flashes when a shock is advised. You must press the button for the shock to be delivered.
shockable rhythm	A heart rhythm that the HeartStart FRx Defibrillator determines is appropriate for defibrillation, such as ventricular fibrillation and some ventricular tachycardias associated with sudden cardiac arrest.

SMART analysis	The proprietary algorithm used by the HeartStart FRx Defibrillator to analyze the patient's heart rhythm and determine whether a shock is advised.
SMART biphasic waveform	The patented, low-energy defibrillation shock waveform used by the HeartStart FRx Defibrillator. It is an impedance-compensated biphasic waveform. It delivers 150 joules, nominal, into a 50 ohm load; when the Infant/Child Key is inserted, it delivers 50 joules, nominal, into a 50 ohm load.
SMART NSA pause	See "NSA pause."
SMART Pads II	The adhesive pads used with the HeartStart FRx Defibrillator to defibrillate patients of any age or weight. The pads are applied to the patient's bare skin and used to detect the patient's heart rhythm and to transfer the defibrillation shock.
standby mode	The operating mode of the HeartStart FRx Defibrillator when a battery has been installed, and the unit is turned off and ready for use when needed. Shown by blinking green READY light.
standard NSA pause	See "NSA pause."
sudden cardiac arrest (SCA)	Sudden cardiac arrest is the abrupt cessation of the heart's normal pumping of blood, frequently caused by an electrical malfunction in the heart. SCA results in a stoppage of blood flow, absent or abnormal breathing, and unconsciousness.
waveform	See "SMART biphasic waveform."

C GLOSSARY OF SYMBOLS/CONTROLS

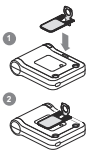
symbol	description
	On/Off button. Green. Pressing the On/Off button when the defibrillator is in standby mode turns the defibrillator on; pressing and holding the On/Off button for one second when the defibrillator is on turns the defibrillator off and disarms the defibrillator. In addition, pressing the On/Off button stops the battery insertion self-test that automatically runs when a battery is inserted.
	Information button (i-button). Pressing the i-button while it is flashing during a patient care pause provides CPR guidance in default configuration; pressing it while it is flashing and the defibrillator is chirping provides troubleshooting guidance. Pressing it until it beeps at other times provides summary information about the defibrillator's last clinical use. Pressing it briefly in standby mode gives device status.
	Caution light. Flashes during rhythm analysis, and is on but not flashing when a shock is advised, as a reminder not to touch the patient.
	Shock button. Orange. If a shock is needed, flashes when the defibrillator is charged. The defibrillator directs the user to press the Shock button to deliver a shock to the patient.
	Consult Instructions for Use. Indicates the need for the user to consult the Instructions For Use. ISO 7000-1641
	Caution. Indicates the need for the user to consult the Instructions For Use for important cautionary information such as warnings and precautions that cannot, for a variety of reasons, be presented on the medical device itself. ISO 7000-0434
	Lithium manganese dioxide battery.
TSO-C142	TSO-C142 certified battery (989803139301 only).

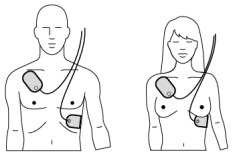
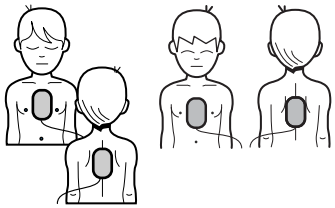
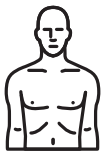
symbol	description
	Environmentally friendly use period (EFUP). Identifies the environmentally friendly use period, up to 50 years.
 QTY (1)	One battery in package.
	Do not charge or recharge the battery.
	Do not deform, damage, or crush the battery.
	Do not expose the battery to high heat, open flames, or dispose of in fire.
	Do not mutilate, dismantle, or open the battery case.
	Needs to be protected from moisture.
	Handle with care.
	This side up.
	Defibrillation-proof type BF applied part. Identifies a defibrillation-proof type BF applied part complying with IEC 60601-1. IEC 60417-5334

symbol	description
IP55	Meets IEC 60529 class IPx5 for jetting water from any direction and class IP5x for protection against access to hazardous parts and ingress of solid foreign objects (dust protected). IEC 60529
C US	Certified by the Canadian Standards Association.
CE	Meets the requirements of the applicable European Directives, including RoHS Directive 2011/65/EU, The Restriction of the Use of Certain Hazardous Substances in Electrical and Electronic Equipment. Directive 93/42/EEC Annex XII
CE 0123	Meets the requirements of the European Medical Device Directive 93/42/EEC. The four numerical digits indicate the identification number of the Notified Body involved in assessing the product's conformity with the directive. Directive 93/42/EEC Annex XII
	Importer. Identifies the entity importing the medical device into the locale.
	European Union. Indicates the European Union.
	Electronic instructions for use. Indicates that relevant information for use of the product is available in electronic form.
	Device Manufacturer.
	Identifies the country of manufacture of products. In the application of this symbol: • The "CC" is replaced by the two-letter or three-letter country code defined in ISO 3166-1. • The date is adjacent to the symbol and in YYYY-MM-DD format.
	Product can be recycled.

symbol	description
	Storage requirements (refer to associated thermometer symbol).
	Transportation requirements (refer to associated thermometer symbol).
	Environmental requirements for transportation and storage.
	Environmental requirements. Indicates the temperature limits to which the medical device can be safely exposed. ISO 7000-0632
	Humidity limitation. Identifies the maximum and minimum limits of relative humidity at which the item shall be stored, transported, or used.
	Atmospheric pressure limit. Indicates the acceptable upper and lower limits of atmospheric pressure for transport and storage.
	Maximum altitude. Identifies that the product is intended to be usable up to the maximum altitude.
	Install-by date. Indicates the date after which the battery is not to be installed. ANSI AAMI ISO 15223-1:2016 Clause 5.1.4
	Catalogue number. Indicates the manufacturer's catalogue number so that the medical device can be identified. ISO 7000-2493
	Authorized representative in the European Community.
	Serial number. Indicates the manufacturer's serial number so that a specific medical device can be identified. ISO 7000-2498
	Batch code. Indicates the manufacturer's batch code so that the batch or lot can be identified. ISO 7000-2492

symbol	description
	Medical device. Identifies the item as a medical device.
	Quantity. Identifies the number of pieces in the package.
	Class 9 miscellaneous dangerous goods. (Symbol required on outer packaging by freight carrier regulations to identify shipments containing lithium batteries.) 49 CFR 172.446
	On HeartStart SMART Pads II (989803139261 only). These pads are disposable and are for single patient use only. ANSI AAMI ISO 15223-1:2016 5.4.2
	Contents: one set of two defibrillation pads.
	Storage temperature limits.
	This product is not made with natural rubber latex.
	This product is not sterile. ANSI AAMI ISO 15223-1:2016 5.2.7
	Replace pads after 24 hours.
	Use-by date. Indicates the date after which the medical device is not to be used. ANSI AAMI ISO 15223-1:2016 5.1.4
YYYY-MM-DD	Expiration date.
	Non-ionizing electromagnetic radiation. Indicates equipment or systems that include RF transmitters or intentionally apply RF electromagnetic energy for diagnosis or treatment.

symbol	description
℞ ONLY Rx only	Federal law (USA) restricts this device to sale by or on the order of a physician. Product or item to be used only per doctor's written prescription. FDA Final Rule - Symbols in Labeling (2016)
	MR Unsafe. Do not use the HeartStart in a magnetic resonance environment. ASTM F2503
	Not for use with Laerdal defibrillator models 911, 1000, 2000, or 3000.
	Not for use with HeartStart HS1 defibrillators, including HeartStart Home and HeartStart OnSite.
	Fits Philips HeartStart designated connector ports, including FRx, FR3, FR2+, and MRx.
< 55 LBS / 25 KG	For use on infants and children under 55 pounds (25 kg).
	Insert Infant/Child Key into slot on FRx.
	Separate collection for Waste Electrical and Electronic Equipment (WEEE) is required. EN 50419:2006 Figure 1
2010 GUIDELINES	Indicates the AHA/ERC/ILCOR resuscitation Guidelines version for which the device is optimized (expressed as a year).
UDI 	Example of the Unique Device Identification (UDI) bar code.

symbol	description
	Proper pads placement for adults.
	Infant/Child with single pad placement illustration. Proper location of pads placement on a pediatric patient.
	Proper location of pads placement on a pediatric patient.
	Pads to be used on adult patients.
	Pads to be used on pediatric patients.

Intentionally blank.

D TECHNICAL INFORMATION

HEARTSTART FRx 861304 DEFIBRILLATOR SPECIFICATIONS

The specifications provided in the following tables are nominal values.

PHYSICAL

category	specifications
size	2.4 in H x 7.1 in D x 8.8 in W (6 cm H x 18 cm D x 22 cm W).
weight	Approximately 3.5 lb (1.6 kg) with battery and pads installed.
pads compatibility	HeartStart SMART Pads II 989803139261
service life	The FRx has an expected service life of 10 years.

ENVIRONMENTAL

category	specifications
temperature and relative humidity	Operating (battery installed, pads connected): 32°F to 122°F (0°C to 50°C); 0% to 95% RH (non-condensing). Standby (between uses with battery installed and pads connected): 32°F to 122°F (0°C to 50°C); 10% to 75% RH (non-condensing). Shipping (with battery and pads not installed): -4°F to 140°F (-20°C to 60°C) for up to 1 week; 0% to 85% RH (non-condensing) for up to 2 days, thereafter 65% RH maximum. Transient Operating (for 20 minutes or less, after rapid transition from 68°F [20°C]): -4°F to 122°F (-20°C to 50°C); Under non-condensing humidity conditions.
altitude	-1312 ft to 15,000 ft (-400 m to 4,572 m).
atmospheric pressure	1060 hPA to 590 hPA.

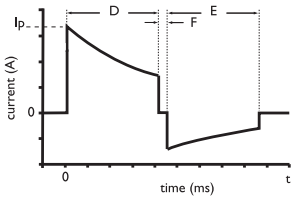
category	specifications
shock/drop abuse tolerance	Withstands 4-foot (1.22 meter) drop on any edge, corner, or face of the device onto masonry surface.
vibration	Operating: meets MILSTD 810G Fig. 5146E-1, random. Standby: meets MILSTD 810G Fig. 5146E-2, swept sine (helicopter).
sealing	Meets IEC 60529 class IP55. Protected against jetting water from any direction per IEC 60529 class IPx5, and against access to hazardous parts and ingress of solid foreign objects (dust protected) per IEC 60529 class IP5x.
crush	1,100 lb (500 kg)
ESD/EMI (radiated and immunity)	See Appendix G
aircraft: method	The FRx meets the following RTCA DO-160G:2010 test levels: Radiated Emissions - Section 21 Category M Conducted Immunity - Section 20 Category M Radiated Immunity - Section 20 Category R and Section 20 Category D Square Wave/Continuous Wave (SW/CW) from 100 MHz to 18 GHz. The FRx has been tested for use in the fuselage of fixed wing aircraft of the following types: Turbojet Engines Reciprocating & Turboprop Engines Reference: RTCA DO-160G Category S, Curves C and L

CONTROLS AND INDICATORS

category	specifications
controls	Green On/Off button i-button (flashes blue) Orange Shock button Optional Infant/Child Key accessory
indicators	Ready light: green, blinks when the defibrillator is in standby mode (ready for use); solid when the defibrillator is being used. i-button: flashes blue when information is available, on solid during patient care pause. Caution light: flashes when the defibrillator is analyzing, comes on solid when the defibrillator is ready to deliver a shock. Shock button: orange, flashes when the defibrillator is charged and ready to deliver a shock. Pads Placement LEDs: flash when FRx is turned on; off once pads are placed on patient. Also operates with Infant/Child Key inserted to indicate pads placement on infants and children under 55 pounds (25 kg) or 0-8 years old.
audio speaker	Provides voice instructions and warning tones during normal use.
beeper	Provides chirps when troubleshooting is needed.
status indicator	Status indicator LCD displays device readiness for use.
low battery detection	Automatic during daily periodic self-testing.
low battery indicator	Alarm chirps and flashing blue i-button.
performance with low battery	When used at temperatures between 50°F and 122°F (10°C and 50°C) with a battery that has just passed the low battery detection threshold, the FRx is capable of delivering a combination of 9 successive therapeutic shocks, separated by no more than 30 seconds and 15 minutes of operation in monitoring mode.

DEFIBRILLATION WAVEFORM

category	nominal specifications
waveform parameters	Biphasic truncated exponential. Waveform parameters are automatically adjusted as a function of patient defibrillation impedance. In the diagram at left, D is the duration of phase 1 and E is the duration of phase 2 of the waveform, F is the interphase delay (500 µs), and Ip is the peak current. The HeartStart FRx delivers shocks to load impedances from 25 to 180 ohms. The duration of each phase of the waveform is dynamically adjusted based on delivered charge, in order to compensate for patient impedance variations, as shown below:



adult defibrillation

load resistance (Ω)	phase 1 duration (ms)	phase 2 duration (ms)	peak current (A)	delivered energy (J)
25	2.8	2.8	55	128
50	4.5	4.5	32	150
75	6.3	5.0	23	155
100	8.0	5.3	18	157
125	9.7	6.4	14	159
150	11.5	7.7	12	160
175	12.0	8.0	11	158

<table>
<tr><th>category</th><th colspan="5">nominal specifications</th></tr>
<tr><td></td><td colspan="5">pediatric defibrillation
(using Infant/Child Key 989803139311)</td></tr>
<tr><td></td><td>load resistance (Ω)</td><td>phase 1 duration (ms)</td><td>phase 2 duration (ms)</td><td>peak current (A)</td><td>delivered energy (J)</td></tr>
<tr><td></td><td>25</td><td>2.8</td><td>2.8</td><td>32</td><td>43.4</td></tr>
<tr><td></td><td>50</td><td>4.5</td><td>4.5</td><td>19</td><td>50.2</td></tr>
<tr><td></td><td>75</td><td>6.3</td><td>5.0</td><td>13</td><td>51.8</td></tr>
<tr><td></td><td>100</td><td>8.0</td><td>5.3</td><td>10</td><td>52.4</td></tr>
<tr><td></td><td>125</td><td>9.0</td><td>6.0</td><td>8</td><td>52.3</td></tr>
<tr><td></td><td>150</td><td>9.0</td><td>6.0</td><td>7</td><td>50.2</td></tr>
<tr><td></td><td>175</td><td>9.0</td><td>6.0</td><td>6</td><td>48.1</td></tr>
<tr><td>energy</td><td colspan="5">Using HeartStart SMART Pads II for adult defibrillation: 150 J nominal (±15%) into a 50 ohm load.

Using HeartStart SMART Pads II with Infant/Child Key inserted: 50 J nominal (±15%) into a 50 ohm load. Sample pediatric energy doses:

age — energy dose
newborn — 14 J/kg
1 year — 5 J/kg
2 - 3 years — 4 J/kg
4 - 5 years — 3 J/kg
6 - 8 years — 2 J/kg

Doses indicated are based on CDC growth charts for the 50th percentile weights.*
* National Center for Health Statistics in collaboration with the National Center for Chronic Disease Prevention and Health Promotion. CDC growth charts: weight-for-age percentiles, modified November 21, 2000. Atlanta, GA: Centers for Disease Control and Prevention © 2000.</td></tr>
<tr><td>charge control</td><td colspan="5">Controlled by Patient Analysis System for automated operation.</td></tr>
<tr><td>shock-to-shock cycle time</td><td colspan="5">< 20 seconds typical, including analysis.</td></tr>
<tr><td>“charge complete” indicator</td><td colspan="5">Shock button flashes, audio tone sounds; device is able to deliver a shock as soon as a shock is advised.</td></tr>
</table>

category	nominal specifications
patient care pause-to-shock time	Quick Shock. 8 seconds, typical, from end of patient care pause to shock delivery.
disarm (AED mode)	Once charged, the HeartStart FRx will disarm if: • patient's heart rhythm changes to non-shockable rhythm, • a shock is not delivered within 30 seconds after the FRx is armed, • the On/Off button is pressed for one second to turn off the FRx, • the Infant/Child Key is inserted or removed, • the battery has been removed or is completely depleted, or • the impedance between pads is out of range.
adult shock delivery vector	Via SMART Pads II placed in the anterior-anterior (Lead II) position.
infant/child shock delivery vector	Via SMART Pads II typically placed in the anterior-posterior position.

ECG ANALYSIS SYSTEM

category	specifications
function	Evaluates impedance of adhesive pads for proper contact with patient skin, and evaluates the ECG rhythm and signal quality to determine if a shock is appropriate.
shockable rhythms	Ventricular fibrillation (VF) and some ventricular tachycardias, including ventricular flutter and polymorphic ventricular tachycardia (VT). The HeartStart FRx Defibrillator uses multiple parameters to determine if a rhythm is shockable. *NOTE: Some very low-amplitude or low-frequency rhythms may not be interpreted as shockable VF rhythms. Also, for safety reasons, some VT rhythms often associated with circulation may not be interpreted as shockable rhythms.*
non-shockable rhythms	On detection of any non-shockable rhythm, prompts user to perform CPR if needed.
pacemaker detection	Pacemaker artifact is removed from the signal for rhythm analysis.
artifact detection	If electrical "noise" (artifact) is detected that interferes with accurate rhythm analysis, analysis will be delayed until the ECG signal is clean.
analysis protocol	Depending on results of analysis, either prepares for shock delivery or provides a pause. For details of protocol, see Appendix E.

ECG ANALYSIS PERFORMANCE

rhythm class	ECG test sample[a] size	meets AHA recommendations[b] for adult defibrillation	
		observed performance	90% one-sided lower confidence limit
shockable rhythm — ventricular fibrillation	300	sensitivity >90%	(87%)
shockable rhythm — ventricular tachycardia	100	sensitivity >75%	(67%)
non-shockable rhythm — normal sinus rhythm	300	specificity >99%	(97%)
non-shockable rhythm — asystole	100	specificity >95%	(92%)
non-shockable rhythm — all other non-shockable rhythms[c]	450	specificity >95%	(88%)

a. From Philips Medical Systems ECG rhythm databases.

b. American Heart Association (AHA) AED Task Force, Subcommittee on AED Safety & Efficacy. Automatic External Defibrillators for Public Access Use: Recommendations for Specifying and Reporting Arrhythmia Analysis Algorithm Performance, Incorporation of New Waveforms, and Enhancing Safety. *Circulation 1997;95:1677-1682*.

c. Supraventricular tachycardia (SVT) is specifically included in the non-shockable rhythm class, in accordance with AHA recommendations[b] and the AAMI standard DF80.

ACCESSORIES SPECIFICATIONS

HEARTSTART SMART PADS II 989803139261

category	specifications		
pads for defibrillation, pacing, monitoring, cardioversion	Disposable, adhesive pads with a nominal active surface area of 12.4 in^2 (80 cm^2) each, provided in a disposable plastic case, and an integrated 48 in (122 cm), typical, cable. Pads in case are designed to fit into carry cases.		
SMART Pads II compatibility	defibrillator model	adult patient use	infant/child patient use
	FRx* FR3 FR2/FR2+ MRx/XL HS1/OnSite/Home	yes yes yes yes no; use M5071A	yes yes no; use M3870A manual mode only no; use M5072A
	* Pre-connectable to FRx defibrillator only.		
pads shelf life	Pads package is labeled with a use-by date of at least two years from date of manufacture.		

M5070A BATTERY AND FRx AVIATION* BATTERY

category	specifications
battery type	9 VDC, 4.2 Ah, lithium manganese dioxide. Disposable, long-life primary cell.
capacity	When new, a minimum of 200 shocks or 4 hours of operating time at 77°F (25°C).
shelf life (prior to insertion)	A minimum of 5 years from date of manufacture when stored and maintained according to directions provided in this document.
standby life (after insertion)	Typically, 4 years when stored and maintained according to directions provided in this document.
training life	Supports 10 hours of use in training mode.

* FRx Aviation Battery Pack (989803139301) complies with TSO-C142. The conditions and tests for TSO approval of this article are minimum performance standards. Those installing this article, on or in a specific type or class of aircraft, must determine that the aircraft installation conditions are within the TSO standards. TSO articles must have separate approval for installation in an aircraft. The article may be installed only according to 14 CFR part 43 or the applicable airworthiness requirements. Lithium cell and battery safety concerns include the possibility of fire and venting of toxic gases.

category	specifications
battery limitations	Never charge, short circuit, puncture, deform, incinerate, heat above 140°F (60°C), or expose contents to water. Remove the battery when discharged.
maintenance and calibration requirements for continued airworthiness (989803139301 only)	There are no periodic maintenance or calibration requirements that are necessary for continued airworthiness of the 989803139301 battery. For battery maintenance with respect to performance within the FRx, please see Chapter 5. There are no user-serviceable parts in the battery.
environmental qualification per RTCA/ DO-227, Section 2.3	Meets following acceptance criteria: No leaking, venting, distortion, fire, or rupture. Change in open circuit voltage <2%.

INFANT/CHILD KEY 989803139311

category	specifications
size	6.3 in x 2.4 in x 0.2 in (16 cm x 6 cm x 0.5 cm).
weight	1.0 oz (29 g).
material	Polycarbonate.

ENVIRONMENTAL CONSIDERATIONS

By complying with your national or local regulations regarding disposal of electric, electronic, and battery waste, you can make a positive contribution to our shared environment.

product	information
defibrillator	The defibrillator contains electronic components. Do not dispose of it as unsorted municipal waste. Collect such electronic waste separately and dispose of it at an appropriate recycling facility according to your country's or local regulations.
battery	The battery cells contain chemicals. The chemistry used in each battery is identified by a symbol on the label; symbols are defined in the defibrillator User's Guide/Instructions for Use/Owner's Manual. Recycle the battery at an appropriate recycling facility.
pads	The used pads may be contaminated with body tissue, fluid, or blood. Dispose of them as infectious waste. Recycle the case at an appropriate recycling facility.

Intentionally blank.

E CONFIGURATION

OVERVIEW

The Philips HeartStart FRx Defibrillator comes with a factory default configuration designed to meet the needs of most users. This configuration can only be changed by using HeartStart Configure software. This software is for use by trained personnel. Information about HeartStart data management products is available online www.philips.com/eventreview. See Appendix A for ordering information.

DEVICE OPTIONS

The following table includes the features of FRx operation that are not related to patient treatment.

parameter	settings	default	default description
speaker volume	1, 2, 3, 4, 5, 6, 7, 8	8	The volume of the FRx speaker is set to 8, highest.
auto send periodic self-test (PST) data	On, Off	On	Enables the periodic self-test data to be broadcast through the device's infrared data port.
ECG out data	On, Off	On	Enables the ECG data to be broadcast through the device's infrared data port.

PATIENT TREATMENT PROTOCOL OPTIONS

parameter	settings	default	default description
"call EMS" voice reminder timing	• At power on (when the user turns on the FRx) • At power on and at the start of the first patient care pause • At the start of the first patient care pause • No reminder	At the start of the first patient care pause.	Provides a voice reminder to make sure emergency medical services have been called, at the start of the first patient care pause.

parameter	settings	default	default description
shock series	1, 2, 3, 4	1	The automatic protocol pause for CPR is activated each time a shock is delivered.* During the protocol pause, the FRx does not perform rhythm analysis. The length of the protocol pause after a shock series is completed is determined by the protocol pause timer setting.
* A shock series begins when a shock is delivered after the FRx is turned on. A new shock series begins after a protocol pause. If shock series is configured for 2 or more, a new shock series also begins if the time since the previous shock exceeds the shock series interval setting.			
shock series interval (minutes)	1.0, 2.0, ∞ (infinity)	1.0	A delivered shock must occur within 1 minute of the previous shock to be counted as part of the current shock series. *NOTE: This parameter is only applicable when the shock series is not configured to the default 1 shock.*
protocol pause timer (minutes)	0.5, 1.0, 1.5, 2.0, 2.5, 3.0	2.0	A 2-minute protocol pause for CPR automatically starts after voice instruction is given when a shock series is completed. After the protocol pause, the defibrillator returns to rhythm analysis. If the user presses the i-button for optional CPR guidance, the FRx provides guidance for 5 cycles of CPR, starting and ending with compressions, when the CPR guidance parameters are also set to their default values. The number of CPR cycles varies for other protocol pause timer and CPR guidance parameter settings.

parameter	settings	default	default description
NSA pause type	• Standard NSA pause: FRx does not perform rhythm analysis during the NSA pause. • SMART NSA pause: FRx conducts background monitoring during the SMART NSA pause. If a potentially shockable rhythm is detected, FRx terminates the SMART NSA pause and resumes rhythm analysis.	SMART NSA pause	During a SMART NSA pause, the defibrillator conducts background monitoring. If a potentially shockable rhythm is detected in a motionless patient, the defibrillator terminates the SMART NSA pause and resumes rhythm analysis. *NOTE: If the FRx detects CPR in progress or if the responder has pressed the i-button for CPR guidance, the SMART NSA pause will be converted to a standard NSA pause. During the standard NSA pause, the defibrillator does not perform rhythm analysis.*
NSA pause timer (minutes)	0.5, 1.0, 1.5, 2.0, 2.5, 3.0	2.0	A 2-minute NSA pause for CPR automatically starts after voice instruction is given when no shock is advised (NSA).* If the user presses the i-button for optional CPR guidance, the FRx provides guidance for 5 cycles of CPR, starting and ending with compressions, when the CPR guidance parameters are also set to their default values. The number of CPR cycles varies for other NSA pause timer and CPR guidance parameter settings.

* If the shock series is configured to 2 or more, and a shock has been delivered as part of a series, the length of the first NSA pause interval within that shock series is determined by the protocol pause timer setting. Otherwise, the length of an NSA pause is determined by the NSA pause timer setting.

parameter	settings	default	default description
CPR prompt	• CPR1: Instructs the user to begin CPR. • CPR2: Instructs the user that it is safe to touch the patient and to begin CPR. • CPR3: Instructs the user to begin CPR and to press the i-button for CPR guidance. • CPR4: Instructs the user that it is safe to touch the patient, to begin CPR, and to press the i-button for CPR guidance.	CPR4: Instructs the user that it is safe to touch the patient, to begin CPR, and to press the i-button for CPR guidance.	The CPR reminder voice instructions provided at the beginning of a pause interval assures the user that it is safe to touch the patient, instructs the user to begin CPR, and invites the user to press the i-button for guidance in the basic steps of CPR. *NOTE: CPR guidance is available only with the CPR3 and CPR4 settings.*
CPR guidance adult ventilation instruction	yes, no	yes	Optional CPR guidance includes rescue breaths at the rate determined by the CPR guidance compression: ventilation ratio for adults when an adult pads set is installed. *NOTE: If this parameter is configured to NO, CPR guidance will always be compressions-only when an adult pads set is installed.*
CPR guidance infant/child ventilation instruction	yes, no	yes	Optional CPR guidance includes rescue breaths at the rate determined by the CPR guidance compression: ventilation ratio for infants and children when an infant/child pads set is installed. *NOTE: If this parameter is configured to NO, CPR guidance will always be compressions-only when an infant/child pads set is installed.*

parameter	settings	default	default description
CPR guidance compression:ventilation ratio	• 30:2 adult and 30:2 infant/child • 30:2 adult and 15:2 infant/child • 15:2 adult and 15:2 infant/child	30:2 adult and 30:2 infant/ child	When the user presses the i-button for optional CPR guidance during a protocol pause or NSA pause, the FRx will provide guidance in basic CPR for cycles of 30 compressions and 2 ventilations for adults, children, and infants. Pauses begin and end with compressions.

Intentionally blank.

F TESTING AND TROUBLESHOOTING

TESTING

The HeartStart FRx Defibrillator automatically tests its battery, connected SMART Pads II, and internal circuitry every day. It alerts you if it finds a problem.

You can also test the defibrillator at any time by removing the battery for five seconds then reinstalling it. This test takes about one minute. Because the battery insertion self-test is very detailed and uses battery power, running it more often than necessary will drain the battery prematurely. It is recommended that you run the battery insertion self-test only:

- when the defibrillator is first put into service.
- after each time the defibrillator is used to treat a patient.
- when the battery is replaced.
- when the defibrillator may have been damaged.

NOTE: If the FRx turns off when you install the battery instead of running the battery insertion self-test, check to be sure that the pads case is not open. If the pads case is open, the FRx assumes that it may be in use and so will not run the self-test.

If you need to use the defibrillator to treat a victim of SCA while you are running a battery self-test, press the On/Off button to stop the test and turn on the HeartStart FRx for use.

TROUBLESHOOTING

The FRx green Ready light is the signal that tells you if the defibrillator is ready for use. The defibrillator chirps and the i-button flashes to alert you to a problem.

RECOMMENDED ACTION WHEN YOU NEED TO USE THE DEVICE

If the FRx is chirping and the i-button is flashing, it is possible that the defibrillator still has enough battery power to be used to treat a victim of SCA. Press the On/Off button.

If the FRx does not turn on when you press the On/Off button, remove the battery and replace it with a new battery if available and press the On/Off button to turn on the defibrillator. If no spare battery is available, remove the installed battery for five seconds, then reinsert it and run a battery insertion self-test.

If the problem continues, do not use the defibrillator. Attend to the patient, providing CPR if needed, until emergency medical personnel arrive.

TROUBLESHOOTING WHILE THE FRx IS BEING USED

(green Ready light is solid)

Always follow any instructions the device gives.

defibrillator says:	possible cause	recommended action
... to replace the battery immediately	The battery is nearly depleted. The FRx will turn off unless a new battery is installed.	Install a new battery immediately.
... to plug in pads connector	The pads connector has been unplugged.	Plug in the pads connector.
... to replace pads	• The pads have been damaged. • The pads have been peeled from the case, but have not been successfully attached to the patient. There may be a problem with the pads.	• Replace the damaged pads. • Replace the pads on patient with new pads to continue with the rescue.
... to press the pads firmly to the skin	The pads are not properly applied to the patient.	Make sure that the pads are sticking completely to the patient's skin.

defibrillator says:	possible cause	recommended action
... to make sure the pads have been removed from the case	The pads are not making good contact with the patient's bare chest because of moisture or excessive hair.	If the pads are not sticking, dry the patient's chest and shave or clip any excessive chest hair.
... the pads should not be touching the patient's clothing	• The pads are touching each other. • The pads may not have been removed from the case or may be on the patient's clothing.	• Reposition the pads. • Make sure pads are not in the case or on patient's clothing.
... to make sure the pads connector is fully inserted	Pads connector is not fully inserted.	Make sure the pads connector is fully inserted.
If the voice instruction continues after you do these things, replace the pads set.		
... to stop all motion	The patient is being moved or jostled.	Stop CPR; do not touch the patient. Minimize patient motion. If the patient is being transported, stop the vehicle.
	The environment is dry and movement around the patient is causing static electricity to interfere with ECG analysis.	Responders and bystanders should minimize motion, particularly in dry environments that can generate static electricity.
	Radio or electrical sources are interfering with ECG analysis.	Check for possible causes of radio and electrical interference and turn them off or remove them from the area.
... the shock was not delivered	The pads may not be making good contact with the patient's skin.	Press the pads firmly to the patient's chest.
	The pads may be touching each other.	Make sure the adhesive pads are correctly positioned on the patient.
	The pads may be damaged.	Replace the pads if necessary.
... the shock button was not pressed	Shock has been advised but the shock button has not been pressed within 30 seconds.	When next prompted, press the Shock button to deliver shock.

TROUBLESHOOTING WHILE THE FRx IS NOT BEING USED

(green Ready light is not on)

Press the i-button to check defibrillator status, and follow any instructions the device gives.

NOTE: In the event of a triple-chirp alert, even if the failure is cleared by a battery insertion test, please contact Philips for service. In the event of repeated instances of a self-test failure resulting in single-chirp alerts, even if such failures are cleared by a battery insertion test, please contact Philips for service.

behavior	possible cause	recommended action
chirps or i-button flashes	The battery power is low or the pads need to be replaced.	Press the flashing blue i-button. Replace the battery or pads if instructed.
	The pads may be damaged or the adhesive dried out.	Replace the pads with a new set and do not open the case until pads are needed in an emergency.
	The pads case may be open.	Make sure the pads case is closed.
	The defibrillator may have been turned off without a pads set installed.	Make sure the pads are properly installed. (See Chapter 2 for directions.)
	The Training Pads II set has been left in the defibrillator.	Remove the Training Pads II set and replace it with a set of SMART Pads II.
	The Infant/Child Key may have been left installed.	Remove the Infant/Child Key.
	The defibrillator has been stored outside the recommended temperature range.	Remove the battery for five seconds then reinstall it to start the battery insertion self-test. If it fails, insert a new battery to repeat the test. If it fails again, do not use the defibrillator. If it passes, store the defibrillator within the recommended temperature range.
	The defibrillator has detected an error during a self-test or cannot perform a self-test, or the Shock button is damaged.	Contact Philips for service.

behavior	possible cause	recommended action
no chirping and/or i-button does not flash, or no response to pressing i-button	The battery is missing or completely depleted.	Remove the battery for five seconds then reinstall it to start the battery insertion self-test. If it fails, insert a new battery and repeat the test. If it fails again, do not use the defibrillator.
	The defibrillator may have been physically damaged.	Contact Philips for service.

Intentionally blank.

G ELECTROMAGNETIC COMPATIBILITY AND IMMUNITY INFORMATION

ELECTROMAGNETIC CONFORMITY

Guidance and manufacturer's declaration: The Philips HeartStart FRx Defibrillator has been tested to demonstrate compliance to the following standards.

COMPLIANCE STANDARDS

IEC standard	edition of compliance and equivalent standards
IEC 60601-1-2	IEC 60601-1-2:2014 EN 60601-1-2:2015 ANSI/AAMI/IEC 60601-1-2:2014 CAN/CSA-C22.2 NO. 60601-1-2:16
CISPR 11	CISPR 11:2009+A1:2010 CISPR 11:2015+AMD1:2016
IEC 61000-4-2	IEC 61000-4-2:2008 EN 61000-4-2:2009 CAN/CSA-IEC 61000-4-2:12 (R2017)
IEC 61000-4-3	IEC 61000-4-3:2006+AMD1:2007+AMD2:2010 EN 61000-4-3:2006+A2:2010 CAN/CSA-CEI/IEC 61000-4-3-07 (R2015)
IEC 61000-4-6	IEC 61000-4-6:2013 EN 61000-4-6:2014 CAN/CSA-IEC 61000-4-6:15
IEC 61000-4-8	IEC 61000-4-8:2009 EN 61000-4-8:2010 CAN/CSA-IEC 61000-4-8:12 (R2017)

Emissions class, group and immunity test levels are shown below for these standards. The FRx is suitable for use in all establishments, including

industrial establishments, domestic establishments, and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes. The customer or user of the FRx should assure that it is used in such an environment.

ELECTROMAGNETIC EMISSIONS

emissions test	compliance	electromagnetic environment – guidance
RF CISPR 11	Group 1 Class B	The FRx 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.
RTCA DO-160	Category M	The FRx may be located in the passenger cabin or cockpit of a transport aircraft.

ELECTROMAGNETIC IMMUNITY

The FRx has been tested according to the levels in RTCA DO-160G Section 20, Category M for conducted immunity. For radiated immunity, the FRx was tested according to Section 20, Category R and Category D Square Wave/Continuous Wave (SW/CW) from 100 MHz to 18 GHz.

Interference may occur in the vicinity of equipment marked with the following symbol:



The HeartStart FRx is not suitable for use next to High Frequency (HF) Surgical Equipment.

Excessive electromagnetic interference may interfere with the FRx's ability to interpret the patient's heart rhythm and/or its ability to deliver therapy.

MRI SAFETY INFORMATION

The HeartStart FRx Defibrillator is MR Unsafe.

ELECTROMAGNETIC IMMUNITY TABLE

The FRx complies with the immunity requirements of IEC 60601-1-2:2014. The immunity test levels for IEC 60601-1-2:2014 are provided in the table below.

immunity test	IEC 60601-1-2:2014 Home Health Environment test level		deviation	notes or special requirements
electrostatic discharge (ESD) IEC 61000-4-2	±8 kV contact*	±2 kV, ±4 kV, ±8 kV, ±15 kV air	Add ±2 kV, ±4 kV, ±6 kV contact	There are no special requirements with respect to electrostatic discharge[a].
power frequency magnetic field IEC 61000-4-8	30 A/m	50 Hz or 60 Hz*	50 Hz and 60 Hz	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial/hospital environment. There are no special requirements for non-commercial/non-hospital environments.
conducted RF IEC 61000-4-6	3 Vrms (outside ISM bands[b])	150 kHz to 80 MHz[d]	N/A	Recommended separation distance[c]: $d = 1.2 \sqrt{P}$
	6 Vrms* (in ISM bands[b])		10 Vrms (in ISM bands)	
radiated RF IEC 61000-4-3	10 V/m*	80 MHz to 800 MHz[d]	20 V/m	$d = 0.60 \sqrt{P}$
		800 MHz to 2.7 GHz		$d = 1.2 \sqrt{P}$

NOTE: For values marked with an asterisk (*), the deviation applies.

a. Generally, AEDs are sometimes susceptible to interference generated by patient and/or responder motion in environments in which a high static electric field is present (e.g., low humidity, synthetic carpets, etc.). As a safety measure, Philips AEDs incorporate a patented method to sense possible corruption of the ECG signal by such interference and to respond by directing the user to stop all motion. In these cases, it is important to minimize movement in the vicinity of the patient during rhythm analysis in order to ensure that the signal being analyzed accurately reflects the patient's underlying heart rhythm.

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

c. The compliance levels in the ISM frequency bands between 150 kHz and 80 MHz and in the frequency range 80 MHz to 2.7 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.

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

Electromagnetic Immunity Table (Continued)

immunity test	IEC 60601-1-2:2014 Home Health Environment test level		deviation	notes or special requirements
RF communications bands specified by Table 9 of IEC 60601-1-2:2014 using IEC 61000-4-3	9 V/m	710 MHz, 745 MHz, 780 MHz, 5240 MHz, 5500 MHz, 5785 MHz	N/A	Recommended separation distance: $d = 0.67 \sqrt{P}$
	27 V/m	385 MHz	N/A	$d = 0.22 \sqrt{P}$
	28 V/m	450 MHz, 810 MHz, 870 MHz, 930 MHz, 1720 MHz, 1845 MHz, 1970 MHz, 2450 MHz		

NOTE 1: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects, and people.

NOTE 2: Recommended separation distance is calculated using the formula shown in the table, where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in meters (m).

NOTE 3: Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, should be less than the compliance level in each frequency range. Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast, and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the FRx is used exceeds the applicable RF compliance level above, the FRx should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such a re-orienting or relocating the FRx.

NOTE 4: The minimum separation distance is 1 foot or 30 centimeters (d = 0.3 m).

RECOMMENDED SEPARATION DISTANCES BETWEEN PORTABLE AND MOBILE RF COMMUNICATIONS EQUIPMENT AND THE HEARTSTART FRx DEFIBRILLATOR

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

rated maximum output power of transmitter (W)	separation distance according to frequency of transmitter[a] (m)		
	150 kHz to 80 MHz[b] $d = 1.2 \sqrt{P}$	80 MHz to 800 MHz[b] $d = 0.60 \sqrt{P}$	800 MHz to 2.7 GHz $d = 1.2\sqrt{P}$
0.01	0.12	0.060	0.12
0.1	0.38	0.19	0.38
1	1.2	0.60	1.2
10	3.8	1.9	3.8
100	12	6.0	12
NOTE 1: For transmitters rated at a maximum output power not listed above, the recommended separation distance *d* in meters (m) can be determined 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.			

a. Excludes the RF communications bands listed in the table on page 74.

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

Intentionally blank.

H SHOCK CYCLE TIMING

The Quick Shock feature allows the FRx to deliver a shock within 8 seconds, typical, following a CPR pause. From shock to shock, the FRx takes <20 seconds, typical, including analysis. After 15 shocks, the FRx takes <30 seconds from analyzing to ready-to-shock. After 200 shocks, the FRx takes <40 seconds from initial power-on to ready-to-shock.

Intentionally blank.

Intentionally blank.

Intentionally blank.

Intentionally blank.

Intentionally blank.

PHILIPS

United States
Philips Medical Systems
22100 Bothell Everett Highway
Bothell, WA 98021-8431, USA
(800) 263-3342

Canada
Philips Canada
1875 Buckhorn Gate
Mississauga, ON
L4W 5N9, Canada
(800) 567-1080



www.philips.com/IFU

MD

Philips Medical Systems
22100 Bothell Everett Highway
Bothell, WA 98021-8431, USA

45356498251 1