

## **Safety Notice**

### **Medical Devices**

# HeartStart FRx, HeartStart HS1 (Home) and HeartStart HS1 (OnSite) AEDs

**Priority 2 – Warning** 





HPRA Safety Notice: SN2018(37) Issue Date: 24<sup>th</sup> October 2018

MANUFACTURER / SUPPLIER	HPRA CASE REFERENCE
Philips Medical Systems	V35048

#### **ISSUE**

Philips Medical Systems has become aware of an issue affecting a resistor component in approximately 660,000 **HeartStart FRx, HeartStart HS1 (Home) and HeartStart HS1 (OnSite) AEDs** manufactured between 2002-2013. Philips Medical Systems has distributed a field safety notice (FSN) to highlight this issue, emphasise the purpose and meaning of audible chirps to users and to notify users of steps to take in the rare event that a shock is not delivered when needed.

Users of devices affected by this notification and whose devices are still under warranty, can receive a replacement AED as detailed in the accompanying FSN.

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#### **ACTION OR RECOMMENDATIONS**

The HPRA advises that users:

- 1 Read the accompanying Field Safety Notice (FSN) carefully.
- 2 Acknowledge receipt of the FSN if you have not already done so.
- Forward a copy of this safety notice and FSN to all relevant personnel within your organisation and to any other organisations or persons where these devices have been transferred.
- 4 Report any concerns regarding these devices and this issue to the manufacturer and the HPRA.
- 5 Contact Philips and/or your distributor to determine whether you are eligible for a replacement device.

#### **TARGET GROUPS**

A&E departments EBME departments

All clinical departments Health and safety managers

All clinical staff
All wards
Ambulance services directors

Maintenance staff
Medical directors
Nursing directors

Ambulance staff Risk managers

Biomedical engineering staff

Volunteer Ambulance Services

Cardiology departments First Aid Organisations

Cardiothoracic departments Sport Centres
Nursing Homes Schools

Work Places/Employers Gyms

Retailers Dental Clinics

#### **BACKGROUND**

**Medical Centres** 

Philips Medical Systems issued this Field Safety Notice (attached) following identification of a resistor component failure in certain AEDs. Typically, AEDs perform various self-tests at daily, weekly and monthly intervals and alert users though audible chirps if an issue is identified. For AEDs effected by this issue, these self-tests are effective at catching the vast majority of issues and alerting users through a series of audible chirps. However, failures have occurred that were not previously detected by these self-tests.

This FSN aims to remind users about the nature and meaning of audible chirps and to provide additional information on the steps to take should audible chirps occur. Further information on these triple chirps and their meaning can be found in the accompanying FSN.

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Users can also view an instructional video available on the Philips website at <a href="https://www.philips.com/aedaudiblechirps">www.philips.com/aedaudiblechirps</a>.

At the time of FSN distribution, Philips Medical Systems was aware of 13 incidents where this component failed during treatment. On these occasions the device delivered at least one shock before failure. Among the cases for which the patient outcome is known; 5 patients died and 2 patients were successfully resuscitated and survived.

The FSN also informs users with affected AEDs of their eligibility for a replacement HeartStart FRx, HeartStart HS1 (Home) and HeartStart HS1 (OnSite) AEDs. Please see the associated FSN for further details. Please contact Philips Medical Systems using the contact details below to determine if you are eligible for a replacement device.

#### **MANUFACTURER**

Enquiries to the **manufacturer** should be addressed to:

 Philips Healthcare UK,
 Telephone:
 00 44 0870 532 9741

 The Philips Centre,
 Fax:
 00 44 01483 369 037

Guildford Business Park, E-mail:

GU2 8XG <u>devicevigilanceuki@philips.com</u>

Website: <u>www.philips.co.uk</u>

#### HPRA CONTACT INFORMATION

All **adverse incidents** relating to a medical device should be reported to:

Health Products Regulatory Authority

Kevin O'Malley House

Fax: +353-1-6764971

Fax: +353-1-6344033

Earlsfort Centre

E-mail: devicesafety@hpra.ie

Earlsfort Terrace Website: www.hpra.ie

Dublin 2

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