



DEFIBRILLATORS RECALLED AFTER REPORTED INCIDENTS

Welch Allyn is recalling about 14,000 external defibrillators after 39 reported incidents, including two that involved patient deaths. The recall, announced March 10, involves 14,054 AED 10 and MRL JumpStart external defibrillators made between October 3, 2002 and January 25, 2007. The Beaverton, Oregon company says there is a remote chance the devices, available through prescription, may produce low-energy shock, shutdown unexpectedly, or be susceptible to electromagnetic noise interference. The issues might prevent defibrillation of a patient in cardiac arrest and could lead to death, the company said in a statement. The company had received 20 instances of low-energy shock, eight instances of electromagnetic noise interference, and 11 instances of the device unexpectedly shutting down.

Welch Allyn AED 10 Recall Information

Important Notice for All Welch Allyn AED 10 Customers

If you purchased a Welch Allyn AED10 or MRL JumpStart defibrillator, your product may be subject to the voluntary recall we initiated on February 25, 2009. Depending upon the date of manufacture, your unit may have a remote chance of having one or more of the following problems:

[Low Energy Shock](#)

[Electromagnetic Noise Interference](#)

[Unexpected Shutdown During Use](#)

[Blown Fuse](#)

[Loss of Voice Prompts](#)

[Shutdown in Cold Temperatures](#)

To determine if your device is subject to this recall, you can use the serial number lookup tool, or you can look for the product's serial number in each of the attached documents below associated with the respective recall issue categories.

http://www.welchallyn.com/support/customer/AED_lookup.jsp



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