

## AED RECALL

Welch Allyn is recalling some of its AEDs. For more information and to determine if your AED is involved, go to [http://www.welchallyn.com/support/customer/AED\\_lookup.jsp](http://www.welchallyn.com/support/customer/AED_lookup.jsp).

The information below is from the FDA's Enforcement Report:

### **PRODUCT**

Welch Allyn AED 10 Automated External Defibrillator. Recall # Z-1005-2009

### **CODE**

Serial numbers: 002664, 002665, 003449, 003875, 003903, 004085, 004108, 004126, 005032, 005096, 005097, 005394, 005438, 006297, 6425, 6728, 6730, 007036, 007039, 007041, 007056, 007057, 007418, 007559, 009501, 010193, 010194, 010196, 010197, 010387, 011026, 011470, 011834, 011936, 012287, 012659, 012859, 013520, 013928, 014655, 014733, 014927, 015154, 015568, 017521, 017683, 018364, 018777, 018941, 019292, 019619, 020422, 020964, 020965, 021543, 021559, 021643, 022509, 022522, 022545, 022546, 022549, 022578, 022579, 022584, 022596, 022597, 022631, 022641, 022654, 023419, 023420, 023421, 023423, 023697, 024000, 025546, 026137, 026878, NA030786, NA031363, NA031533, NA031857, NA032151, NA032179, and NA035807

### **RECALLING FIRM/MANUFACTURER**

Recalling Firm: Welch Allyn Protocol, Inc., Beaverton, OR, by a Service Bulletin on April 13, 2007. Manufacturer: MRL, Inc., a Welch Allyn Company, Buffalo Grove, IL. Firm initiated recall is ongoing.

### **REASON**

Potential for device to shut down prematurely under certain circumstances.

### **VOLUME OF PRODUCT IN COMMERCE**

86 devices

### **DISTRIBUTION**

Nationwide and countries of Belgium, Canada, France, Ireland, Israel, Mexico, and United Kingdom

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### **PRODUCT**

Welch Allyn AED 10 automatic external defibrillator and MRL Jumpstart (collectively "AED 10"). Recall # Z-1006-2009

### **CODE**

Software version 2.02 or lower. The ECG analysis/noise issue involves all units manufactured before January 2006 when a design/component change was made.

### **RECALLING FIRM/MANUFACTURER**

Welch Allyn Protocol, Inc., Beaverton, OR, by letter dated February 26, 2009. Firm initiated recall is ongoing.

### **REASON**

Reliability issues - potential to shock a non shockable rhythm or not shocking a shockable rhythm;

### **VOLUME OF PRODUCT IN COMMERCE**

10,735 units

### **DISTRIBUTION**

Nationwide and Internationally