



## County of Erie

CHRIS COLLINS  
COUNTY EXECUTIVE

### DEPARTMENT OF HEALTH

ANTHONY J. BILLITTIER IV, M.D., FACEP  
COMMISSIONER OF HEALTH

**TO:** All E.M.S. Providers

**FROM:** Daniel Neaverth, Deputy Commissioner / E.M.S.  
Erie County Division of Emergency Medical Services

**DATE:** February 10, 2010

**SUBJECT:** Cardiac Science Automated External Defibrillator (AED) Recall

Cardiac Science Corporation and the FDA notified healthcare professionals and consumers of a recall of certain Automated External Defibrillators (AEDs). The press release is available on the Food and Drug Administration website at:

<http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm200138.htm> .

This is a new recall, separate from the one distributed on November 16, 2009.

These AEDs were manufactured in a way that makes them potentially susceptible to failure under certain conditions. The units that have been identified were manufactured or serviced between October 19, 2009 and January 15, 2010 and include the following models - Powerheart 9300A, 9300E, 9300P, 9390A, 9390E, CardioVive 92532 and CardioLife 9200G and 9231.

To determine if you have a unit affected by this notice, go to [www.cardiacscience.com/AED195](http://www.cardiacscience.com/AED195) . Cardiac Science offers a tool using the serial number to determine if a unit is involved in the corrective action. At this time, ***Cardiac Science is recommending that the unit be removed from service immediately if it is included in this recall.***

If you have or know any organization that has one of these units, refer them to the FDA notice listed above. This website will provide more detail on the units and corrective action proposed for the AEDs.

If you should have any questions, please contact us at the E.M.S. office at 681-6070.

Cc Anthony Billittier, IV, MD, Commissioner  
Gregory Skibitsky, Commissioner