



Inland Counties Emergency Medical Agency

Serving San Bernardino, Inyo, and Mono Counties

*Virginia Hastings, Executive Director
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DATE: September 18, 2009

TO: ICEMA Prehospital Providers
ICEMA 9-1-1 Receiving Hospitals

FROM: Virginia Hastings
ICEMA Executive Director

**SUBJECT: RECALL NOTICE-LIFEPAK CR Plus Automated External Defibrillators
(Physio-Control, Inc)**

FDA notified healthcare professionals of a Class I recall of certain LIFEPAK CR Plus Automated External Defibrillators (AED) manufactured and distributed from July 9, 2008 through August 19, 2008. An extremely humid environment may cause the affected devices to improperly analyze the heart rhythm and may cause the device to delay or fail to deliver therapy.

Any adverse events or quality problems that may be related to the use of this product should be reported to the FDA's [MedWatch Adverse Event Reporting program online](#), by phone [1-800-332-1088], or by returning the postage-paid [FDA Form 3500](#) by mail or fax [1-800-FDA-0178].

Read the complete MedWatch 2009 Safety summary, including a link to the Class 1 recall notice, at:

<http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm182496.htm>

You are encouraged to report all serious adverse events and product quality problems to FDA MedWatch at www.fda.gov/medwatch/report.htm

VH/mae