



California Medical Device Recall Information



Recall Name

HeartSync Recalls Multi-function Defibrillation Electrodes Due to a Connection Issue

Recall Date	Product Description	Recalling Firm	Recall Reason
11/11/14	Multi-function Defibrillation Electrodes	HeartSync, Inc. Ann Arbor, MI	<i>Due to a connector compatibility issue with Philips FR3 and FRx Defibrillator Units.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	<p>All Lot numbers for the following:</p> <ul style="list-style-type: none"> Catalog Number C100-PHILIPS - Adult Radiotransparent Electrode Catalog Number C100AC-PHILIPS - Adult Radiotransparent Electrode Catalog Number T100LO-PHILIPS - Adult Radiotransparent Electrode Catalog Number T100-PHILIPS Radiotranslucent Electrode Catalog Number T100AC-PHILIPS Radiotranslucent Electrode 	CA , nationwide	Distributed from October 26, 2011 through November 26, 2014.

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm425536.htm>