



California Medical Device Recall Information



Recall Name

HeartSine Technologies Recalls Samaritan Defibrillator Due to Possible Malfunction

Recall Date	Product Description	Recalling Firm	Recall Reason
9/11/12	HeartSine 300/300P Samaritan Public Access Defibrillator (PAD)	HeartSine Technologies, Ltd. Belfast, Northern Ireland	<i>Potential failure to perform as intended due to:</i> a) <i>Prematurely depleted batteries;</i> <i>(and/or)</i> b) <i>Faulty battery management software.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	HeartSine 300/300P PAD Affected unit Serial Numbers: • 0400000501 to 0700032917; • 08A00035000 to 10A0070753; • 10C00200000 to 10C00210106	CA , globally	From 8/01/2004 to 01/31/2011

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm328604.htm>