



California Medical Device Recall Information



Recall Name

St. Jude Medical Updates Recall of Optisure Dual Coil Defibrillation Leads Due to Possible Damage

Recall Date	Product Description	Recalling Firm	Recall Reason
Initial: 11/03/15 Updated: 01/22/16	Optisure Dual Coil Defibrillation Leads Model numbers: <ul style="list-style-type: none"> • LDA220 • LDA220Q • LDA230Q • LDP220Q 	St. Jude Medical, Inc. Sylmar, CA	<i>Potential for leads to be compromised during manufacturing process, which could result in the inability of the defibrillator to deliver electrical therapy to the patient.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	List of affected devices	CA, nationwide	Distribution dates: April 9, 2014 to October 20, 2015

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm483428.htm>