



# California Medical Device Recall Information



## Recall Name

### St. Jude Medical Recalls Optisure Dual Coil Defibrillation Leads Due to Possible Damage

| Recall Date  | Product Description  | Recalling Firm                              | Recall Reason   |
|--------------|--|---|---|
| 11/03/15     | <b>Optisure Dual Coil Defibrillation Leads</b><br><br>Model numbers: <ul style="list-style-type: none"> <li>• LDA220</li> <li>• LDA220Q</li> <li>• LDA230Q</li> <li>• LDP220Q</li> </ul> | <b>St. Jude Medical, Inc.</b><br>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            | <a href="#">List of affected devices</a>   | CA, nationwide                              | Distribution dates:<br><br>April 9, 2014 to<br>October 20, 2015   |

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm483068.htm>