



California Medical Device Recall Information



Recall Name

Cordis Corporation Recalls Optease Retrievable Inferior Vena Cava (IVC) Filter Due to Potential for Incorrect Deployment

Recall Date	Product Description	Recalling Firm	Recall Reason
03/29/13	Optease Retrievable Inferior Vena Cava Filter	Cordis Corporation Miami Lakes, FL	<i>Update to device labeling to minimize the likelihood of implanting the filter backwards.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	All products shipped during the affected time period. An "URGENT Medical Device Correction" letter was issued to customers 04/02/13.	CA , nationwide	Distributed in US: 05/06/2010 to 04/02/2013

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm364364.htm>