



# California Medical Device Recall Information



## Recall Name

**Medtronic Neurosurgery Recalls the  
Duet External Drainage and Monitoring System  
Due to a Potential Defect**

Recall Date	Product Description	Recalling Firm	Recall Reason
6/09/14	Medtronic Duet External Drainage and Monitoring System (EDMS)	<b>Medtronic Neurosurgery, Inc.</b> Goleta, CA	<i>Potential for the patient line tubing to separate from the patient line connectors.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	<ul style="list-style-type: none"> <li>• Interlink Injection Sites, Catalog Number <b>46913</b></li> <li>• SmartSite Injection Sites, Catalog Number <b>46914</b></li> <li>• Interlink Injection Sites, Ventricular, Catheter, Catalog Number <b>46915</b></li> <li>• SmartSite Injection Sites, Ventricular Catheter, Catalog Number <b>46916</b></li> <li>• Interlink Injection Sites, Lumbar Catheter, Catalog Number <b>46917</b></li> </ul> <p><a href="#">List of Affected Lot Numbers</a></p>	<b>CA</b> , nationwide	Distributed from:  April 10, 2013 to May 19, 2014

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm403857.htm>