



# California Medical Device Recall Information



## Recall Name

### Medtronic Recalls Sutureless Connector Intrathecal Catheters Due to Catheter Occlusion

Recall Date	Product Description	Recalling Firm	Recall Reason
06/03/13	Sutureless Connector Intrathecal Catheters  [used with the SynchroMed Implantable Infusion System]	<b>Medtronic, Inc.</b> Minneapolis, MN	<i>There is potential for catheter occlusion and interruption of drug delivery.</i>  <i>The Sutureless Connector has been redesigned to reduce the potential for occlusion at the point where the catheter connects to the implantable drug pump.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	Sutureless Connector Intrathecal Catheters  Model Numbers: <ul style="list-style-type: none"> <li>• 8709SC</li> <li>• 8731SC</li> </ul> Sutureless Revision Kits  Model Numbers: <ul style="list-style-type: none"> <li>• 8596SC</li> <li>• 8578</li> </ul> <i>Use by Date = Aug 14, 2014 or sooner.</i>	<b>CA</b> , nationwide	Product manufacture dates: November 21, 2006 - August 24, 2012.  Distribution dates: January 23, 2007 - March 27, 2013.

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm359126.htm>