



California Medical Device Recall Information



Recall Name

Medtronic Recalls Guidewires Due to Potential for Coating to Detach

Recall Date	Product Description	Recalling Firm	Recall Reason
10/21/13	<u>Product Lines</u> <ul style="list-style-type: none"> • Cougar Nitinol Workhorse Guidewire • Cougar Steerable Guidewire • Zinger Stainless Steel Workhorse Guidewire • Zinger Steerable Guidewire • Thunder Extra-Support Guidewire • Thunder Steerable Guidewire • ProVia Crossing Guidewire • Attain Hybrid Guidewire 	Medtronic, Inc. Danvers, MA	<i>Potential for coating on guidewire surface to delaminate and detach.</i> <i>Detachment of the coating could lead to vascular thrombosis or occlusion.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	<u>List of Suspect Models and Lots</u>	CA, nationwide	Manufactured: April through Sept. 2013 Distributed: May through Oct. 2013

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/Recalls/ucm375363.htm>