



California Medical Device Recall Information



Recall Name

**Teleflex Incorporated Recalls ARROW® International
Intra-Aortic Balloon Catheter Kits
Due to Potential for Separation**

Recall Date	Product Description	Recalling Firm	Recall Reason
02/11/16	ARROW® International <ul style="list-style-type: none"> Intra-Aortic Balloon Catheter Kits Percutaneous Insertion Kits 	Teleflex Incorporated Wayne, PA	<i>The sheath body may become separated from the sheath hub. If the separation occurs, the patient may bleed from the sheath.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	Click here to go to a list of affected Product Codes and Lots Numbers	CA , nationwide	Manufactured: December 1, 2013 to December 1, 2015 Distributed: January 1, 2014 to February 1, 2016

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm491270.htm>