



# California Medical Device Recall Information



## Recall Name

**Hospira Recalls Abbott Acclaim and Hospira Acclaim Encore Infusion Pumps  
Due to Broken Door Assemblies**

Recall Date	Product Description	Recalling Firm	Recall Reason
10/31/13	<ul style="list-style-type: none"> <li>• <b>Abbott Acclaim</b> Infusion Pumps</li> <li>• <b>Hospira Acclaim Encore</b> Infusion Pumps</li> </ul>	<b>Hospira, Inc.</b> Lake Forest, IL	<p><i>Customer reports of broken door assemblies on the infusion pumps.</i></p> <p><i>If the door assembly breaks, it may result in an over-infusion or a delay of therapy.</i></p>
Recall Class	Product Identification	Distribution	Affected Dates
I	<p><b>All Infusion Pumps</b></p> <p><b>List Numbers:</b></p> <p><b>12032</b></p> <p><b>12237</b></p>	<b>CA, nationwide</b>	<p><b>Abbott Acclaim pumps:</b></p> <p>Manufactured <b>Feb 1998 - Nov 1998</b>; Distributed <b>Sept 1998 - Feb 2004</b>.</p> <p><b>Hospira Acclaim Encore pumps:</b></p> <p>Manufactured <b>Feb 1997 to Feb 2010</b>; Distributed <b>Jul 1999 to Nov 2013</b>.</p>

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm395845.htm>