



# California Medical Device Recall Information



## Recall Name

### Covidien Recalls Trellis 6 and Trellis 8 Peripheral Infusion Systems Due to Mislabeled Inflation Ports

Recall Date	Product Description	Recalling Firm	Recall Reason
12/10/14	Peripheral Infusion Systems: <ul style="list-style-type: none"> <li>Trellis 6</li> <li>Trellis 8</li> </ul>	<b>Covidien</b> Plymouth, MN	<p><i>A manufacturing error caused the balloon inflation ports to be mislabeled. This may cause the physician using the device to deflate the balloons in the incorrect order.</i></p> <p><i>If this happens, there is a potential for blood clots to dislodge and move into the lungs.</i></p>
Recall Class	Product Identification	Distribution	Affected Dates
I	Follow the link below for a list of affected models and lot numbers:  <a href="#">Affected Devices</a>	<b>CA</b> , nationwide	U.S. Distribution Dates:  June 6, 2014 to November 13, 2014

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm433765.htm>