



# California Medical Device Recall Information



## Recall Name

### Halyard Health Recalls KimVent\* Microcuff\* Subglottic Suctioning Endotracheal Tubes Due to Component Detachment

Recall Date	Product Description	Recalling Firm	Recall Reason
11/17/14	KimVent* Microcuff* Subglottic Suctioning Endotracheal Tubes	Halyard Health Roswell, GA	<p><i>Complaints received concerning detachment of the cuff inflation line from the tube during use.</i></p> <p><i>This may lead to an air leak between the cuff and the tracheal wall, which may reduce the amount of air that reaches the lungs.</i></p>
Recall Class	Product Identification	Distribution	Affected Dates
I	<p>Click on the links below for affected Product Codes and Lot Numbers:</p> <ul style="list-style-type: none"> <li>• <a href="#">7.0 mm</a></li> <li>• <a href="#">7.5 mm</a></li> <li>• <a href="#">8.0 mm</a></li> <li>• <a href="#">8.5 mm</a></li> <li>• <a href="#">9.0 mm</a></li> </ul>	CA, nationwide	<p><u>Manufacture Dates:</u> November 15, 2013 to October 21, 2014</p> <p><u>Distribution Dates:</u> December 20, 2013 to October 30, 2014</p>

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm428617.htm>