



California Medical Device Recall Information



Recall Name

**Teleflex Medical Recalls
Lifesaver Single Patient Use Manual Resuscitator
Due to Blocked Intake Port**

Recall Date	Product Description	Recalling Firm	Recall Reason
5/15/15	Lifesaver Single Patient Use Manual Resuscitator	Teleflex Medical Research Triangle Park, NC	<i>The oxygen intake port may be blocked which can prevent the bag from filling.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	Follow the link below for: Affected Product Codes and Lot Numbers	CA , nationwide	Manufacturing and Distribution Dates: June 2014 to April 2015

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm453549.htm>