California Medical Device Recall Information

Recall Name
Teleflex Medical Recalls the **Hudson RCI Sheridan Sher-I-Bronch Endobronchial Tube** **Due to Possible Breakage or Separation of the Connector on the Tube**

Recall Date	Product Description	Recalling Firm	Recall Reason
6/01/15	Hudson RCI Sheridan Sher-I-Bronch Endobronchial Tube	**Teleflex Medical,** Research Triangle Park, NC	*Potential for the endobronchial tube's double swivel connector to crack or separate.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	Follow the links below for the affected lot numbers: • Right-Sided Endobronchial Tube • Left-Sided Endobronchial Tube • Double Swivel • Accessory Pack	**CA**, nationwide	Manufacturing and distribution dates: October 2013 to May 2015

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm459801.htm