



California Medical Device Recall Information



Recall Name

OriGen Biomedical Recalls OriGen VV13F Reinforced Dual Lumen ECMO Catheters Due to Potential for Separation

Recall Date	Product Description	Recalling Firm	Recall Reason
3/30/15	OriGen VV13F Reinforced Dual Lumen ECMO Catheters	OriGen Biomedical Austin, TX	<i>Potential for a separation of the clear extension tube from the hub that it is inserted in, which potentially could result in required intervention to prevent permanent impairment/damage.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	Lot: N18549 Expiration: 09/2018	CA , nationwide	Manufactured: September 22, 2014 Distributed: February 16 to March 26, 2015

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm443009.htm>