California Medical Device Recall Information

Recall Name
Vascular Solutions Recalls Langston V2 Dual Lumen Catheters (Models 5540 and 5550) Due to Possible Separation

Recall Date	Product Description	Recalling Firm	Recall Reason
05/27/14	Langston V2 Dual Lumen Catheters **Models: 5540, 5550**	**Vascular Solutions, Inc.** Minneapolis, MN	*Due to possible separation of the inner catheter from the device hub during use.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	**Affected Lot numbers:** 569436 570505 570928 569437 570506 570929 569962 570507 571099 569963 570592 571100 569964 570593 571101 569965 570674 571102 570177 570675 571103 570178 570744 571350 570337 570745 571412 570338 570746 571413 570339 570836 571414 570340 570926 571415 570341 570927 571727 570504	**CA**, nationwide	Distributed from: March 2014 to May 2014

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm404054.htm