California Medical Device Recall Information

Recall Name
Boston Scientific Recalls the Chariot Guiding Sheath Due to the Risk of Shaft Separation

Recall Date	Product Description	Recalling Firm	Recall Reason
11/19/15	Chariot Guiding Sheath	**Boston Scientific Corp.** Marlborough, MA	*Potential for the Chariot Guiding Sheath to become separated from the shaft.*

Recall Class	Product Identification	Distribution	Affected Dates
I	**All Universal Product Numbers (UPNs)**: H74939277645110, H74939277745110, H74939277845110, H74939277645220, H74939277765120, H74939277690210, H74939277665110, H74939277790110, H74939277645120, H74939277865120, H74939277690220, H74939277690110, H74939277865110, H74939277745120, H74939277890120, H74939277845210, H74939277765110, H74939277545210, H74939277790210, H74939277845120, H74939277745210, H74939277545110, H74939277790220, H74939277790120, H74939277745220, H74939277645210, H74939277690120, H74939277665120, H74939277890110	**CA**, nationwide	Manufacturing dates: Mar 17, 2015 to Nov. 10, 2015 Distribution dates: Jun 6, 2015 to Nov. 17, 2015

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm476781.htm