California Medical Device Recall Information

Recall Name
Stryker Recalls Angiodynamics Soft Vu Omni Flush Angiographic Catheter Due to Potential Tip Separation

Recall Date	Product Description	Recalling Firm	Recall Reason
06/01/16	Angiodynamics Soft Vu Omni Flush Angiographic Catheters Model Number: **10732203**	**Stryker Corporation** Tempe, AZ	*Potential that the tip of the catheter may separate from the main body.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	Suspect Lot Numbers: **8248U 119965U** **158623U 175183U** **179688U 191185U** **210813U 225911U** **225918U 225919U** **253202U 277726U** **302844U 309329U** **325680U 343908U** **352756U 352761U** **359352U 363987U** **366572U 369818U** **375883U 772102U** **773196U 773216U** **775985U 7710750U**	**CA**, nationwide	Manufactured between: November 7, 2003 and October 18, 2008

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm512633.htm