State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

Cordis Recalls SUPER TORQUE MB Angiographic Catheter with Radiopaque Marker Bands Due to Potential for Marker Bands to Move or Dislodge

Recall Date	Product Description	Recalling Firm	Recall Reason
07/21/2021	**Cordis SUPER TORQUE MB Angiographic Catheter 5F PIG, 598A,B,C**	**Cordis Corporation,** Miami Lakes, Florida	Usage can entrap the catheter between endovascular device and marker bands, causing bands to move or dislodge

Recall Class	Product Identification	Distribution	Affected Dates
I	**Cordis SUPER TORQUE MB Angiographic Catheter 598A** A polycarbonate hub with braided polyurethane body with 10 gold alloy marker bands and unbraided pigtail	40 units within California	January 2020 – July 2021
I	**Cordis SUPER TORQUE MB Angiographic Catheter 598B** Similar to A with 20 gold alloy marker bands	510 units within California	January 2020 – July 2021
I	**Cordis SUPER TORQUE MB Angiographic Catheter 598C** Similar to A with 20 gold alloy marker bands	415 units within California	January 2020 – July 2021

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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