State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

Tornier Recalls Stryker Tornier Augmented Glenoid Lateralized Baseplate For Manufacturing With 6mm Spacers

Recall Date	Product Description	Recalling Firm	Recall Reason
10/25/2023	**Tornier** stryker Tornier Perform Reversed Augmented Glenoid Lateralized Baseplate, 25mm, Offset +3mm, REF DWJ502	**TORNIER,INC** Bloomington, Minnesota	One lot of Tournier Perform Reversed Lateralized Baseplates were found to have been manufactured with 6mm spacers

Recall Class	Product Identification	Distribution	Affected Dates
II	**Tornier** UDI/DI 10846832062017, Lot Number 1756123	30 Units Nationwide including California	September 2023 and prior

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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