State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

Steris Corporation Recalls Dual Articulating Headrest For Failure To Properly Lock In Place

Recall Date	Product Description	Recalling Firm	Recall Reason
2/7/2024	**Dual Articulating Headrest** Velcro P/N P141210813, REF BF753	**STERIS CORPORATION** Mentor, Ohio	Dual-articulating head rest may not stay in place or may fail to lock into position during a patient procedure which could result in an injury to the patient or a procedural delay.

Recall Class	Product Identification	Distribution	Affected Dates
II	**Dual Articulating Headrest** UDI-DI: 00724995195885	15 Units in California	August through October, 2023

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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