State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

Med-Mizer recalls MR600 and STS500 Patient Lifts

Recall Date	Product Description	Recalling Firm	Recall Reason
3/8/2023	**Med-Riser MR600** Patient Lift	**Med-Mizer Inc.** Batesville, Indiana	Risk of boom pivot failing due to the nut securing the boom becoming loose and the bolt to be unsecure to the flange on the side.
3/8/2023	**Sit to Stand STS500** Patient Lift	**Med-Mizer Inc.** Batesville, Indiana	Risk of boom pivot failing

Recall Class	Product Identification	Distribution	Affected Dates
II	**Med-Riser MR600** UDI-DI: 00852195007308; Serial No.: MR600000 - MR600240	8 Units in California	February 2023 and prior
II	**Sit to Stand STS500** UDI-DI: 00852195007353; Serial No.: STS500000 - STS500036	1 Unit in California	February 2023 and prior

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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