State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

ARJOHUNTLEIGH POLSKA Sp. Z.O.O. Recalls Arjo Medical Beds for Faulty Bed Wheels

Recall Date	Product Description	Recalling Firm	Recall Reason
12/22/2023	**Arjo Medical Beds** Models Enterprise 5000X, Enterprise 8000X, Enterprise 9000X and Citadel, assembled with IndiGo Drive Assistance module and retrofitted with IndiGo Drive Assistance modules	**ARJOHUNTLEIGH POLSKA SP. Z.O.O.** Komorniki, Poland	This product is being recalled due to unintended movement of the bed wheels.
12/22/2023	**Arjo Medical Beds** Models Enterprise 5000X, Enterprise 8000X, Enterprise 9000X and Citadel, assembled with IndiGo Drive Assistance module and retrofitted with IndiGo Drive Assistance modules	**ARJOHUNTLEIGH POLSKA Sp. z.o.o.** Komorniki,Poland	This product is being recalled due to unintended movement of the bed wheels.

Recall Class	Product Identification	Distribution	Affected Dates
II	**Arjo Medical Beds** UDI/DI 0505609734548	83 Units Nationwide including California	December, 2023 and prior
II	**Arjo Medical Beds** UDI/DI 050560973512	846 Units Nationwide including California	December, 2023 and prior

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE Arjo Medical Bed 0505609734548, Arjo Medical Bed 050560973512