

## State of California—Health and Human Services Agency

## California Department of Public Health



Food and Drug Branch - Device Recalls

## CALIFORNIA DEVICE RECALL INFORMATION SHEET

Baxter Recalls Hillrom Centrella Hospital Bed and Accessories for Radiofrequency Emission Interference with other Medical Devices

Recall Date	Product Description	Recalling Firm	Recall Reason
11/9/2022	Hillrom Centrella Hospital Bed with WatchCare Incontinence Management System Product code/Part numbers: P7900B100193 - P7900B300453	Baxter Healthcare Corporation Deerfield, Illinois	Radio frequency emission from a functionning WatchCare device may potentially impact other devices (including, but not limited to telemetry devices, bladder scanner, fetal monitor/doppler, infusion pumps, and insulin pumps).
11/9/2022	Hillrom Hospital Bed Access- ory, Watch Care Incontinence Management System for Centrella Bed Product Code/Part Number P00697905	Baxter Healthcare Corporation Deerfield, Illinois	Radio frequency emission
11/9/2022	Hillrom Hospital Bed Access- ory, Watch Care Incontinence Management System for Progr- essa Bed Product Code/Part Number P00697903	Baxter Healthcare Corporation Deerfield, Illinois	Radio frequency emission
11/9/2022	Hillrom Hospital Bed Accessory, WatchCare Incon- tinence Management System for VersaCare Bed Rev. K Product Code/Part Number P00697902	Baxter Healthcare Corporation Deerfield, Illinois	Radio frequency emission

11/9/2022	Hillrom Hospital Bed Accessory, Watch Care Incontinence Management System for VersaCare Bed Rev. A-J Product Code/Part Number P00697901	Baxter Healthcare Corporation Deerfield, Illinois	Radio frequency emission
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Recall Class	Product Identification	Distribution	Affected Dates
I	Hillrom Centrella Hospital Bed with WatchCare UDI/DI 887761985162, all serial numbers	6407 Devices Nationwide	September 2022 and prior
I	Accessory for Centrella Bed UDI/DI 887761984998, all serial numbers	814 Devices Nationwide	September 2022 and prior
I	Accessory for Progressa Bed UDI/DI 00887761998889, all serial numbers	1137 Devices Nationwide	September 2022 and prior
I	Accessory for VersaCare K UDI/DI 887761998872, all serial numbers	63 Devices Nationwide	September 2022 and prior
I	Accesory for VersaCare A-J UDI/DI 00887761998865, all serial numbers	129 Devices Nationwide	September 2022 and prior

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

