California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Baxter Healthcare Corporation Progressa Bed Surfaces

Recall Date	Product Description	Recalling Firm	Recall Reason
03/26/2025	Progressa Bed Surfaces, intended to be used to treat or prevent pulmonary or other complications associated with immobility Product Codes: P7520A19, P7520A20S, P7520A21, P7520A22S, P7520A23, P7520A24S, P7520A3, P7520A31, P7520A37, P7520A39, P7520A45, and P7520A4S	Baxter Healthcare Corporation	The air bladders inside the mattress may move out of position when the head of the bed is elevated, causing a dip in the mattress.

Recall Class	Product Identification	Distribution	Affected Dates
II	Product Code: P7520A19 UDI-DI: 00887761999893 Product Code: P7520A20S UDI-DI: 00887761999886 Product Code: P7520A21 UDI-DI: 00887761999879 Product Code: P7520A22S UDI-DI: 00887761999862	98 units in California	February and prior

	Product Code: P7520A23 UDI-DI: 00887761999855 Product Code: P7520A24S UDI-DI: 00887761999848 Product Code: P7520A3 UDI-DI: 00887761999770 Product Code: P7520A31 UDI-DI: 00887761999756 Product Code: P7520A37 UDI-DI: 00887761999718 Product Code: P7520A39 UDI-DI: 00887761999701 Product Code: P7520A45 UDI-DI: 00887761999671 Product Code: P7520A4S UDI-DI: 00887761999657 Serial Numbers: A008MW1489		

For additional information, please visit the FDA Website.