California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Baxter Healthcare Corporation Recalls Baxter MiniCap Extended Life PD Transfer Set with Twist Clamp for Potential Rick of Exposure to non-dioxin-like (NDL) polychlorinated biphenyl acids (PCBAs) and NDL polychlorinated biphenyls (PCBs)

Recall Date	Product Description	Recalling Firm	Recall Reason
11/20/2024	Baxter MiniCap Extended Life PD Transfer Set with Twist Clamp, Part Number 5C4482; use in Peritoneal Dialysis	Baxter Healthcare Corporation	Baxter is aware of several recalls by other manufacturers related to the potential risk of exposure to non-dioxin-like (NDL) polychlorinated biphenyl acids (PCBAs) and NDL polychlorinated biphenyls (PCBs) when using certain peritoneal dialysis and hemodialysis devices.

Recall Class	Product Identification	Distribution	Affected Dates
I	UDI/DI 00085412007731, Lot/Serial Numbers: All lots including and manufactured after H19I26088	2,706,558 units Worldwide Distribution (US Nationwide distribution)	November 2024 and prior

For additional information, please visit the FDA Website.