California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

MEDLINE INDUSTRIES, LP – Northfield Recalls Medline convenience kits

Recall Date	Product Description	Recalling Firm	Recall Reason
07/09/2025	Medline convenience kits containing BD Insyte Autoguard Shielded IV Catheters 22 GA x 1.00 IN. labeled as: 1) CC DRAWER 4 IV CIRCULATION, REF ACC010254B; 2) CC ADULT DRAWER 2, REF ACC010256; 3) PEDI EMERGENCY TRAY-BLUE, REF ACC010316A; 4) ADULT DRAWER 4, REF ACC010389C; 5) DRAWER 4 - SUMMIT, REF ACC010627; 6) DRAWER 3, REF ACC010898. .	MEDLINE INDUSTRIES, LP - Northfield	Medline Industries, LP. is issuing a recall for specific Medline kits containing BD Insyte Autoguard Shielded IV Catheters 22 GA x 1.00 IN which was identified that during insertion of the Catheter, the needle is slow to retract or fails to retract.

Recall Class	Product Identification	Distribution	Affected Dates
II	1) REF ACC010254B: UDI/DI 10193489651683 (each), 40193489651684 (case), Lot Number: 24LDA269; 2) REF ACC010256: UDI/DI 10884389559529 (each), 40884389559520 (case), Lot Number: 24LDA106; 3) REF ACC010316A: UDI/DI 10889942831746 (each), 40889942831747 (case), Lot Number: 24LDA823; 4) REF ACC010389C: UDI/DI 10193489472493 (each), 40193489472494 (case), Lot Number: 24LDB092; 5) REF ACC010627: UDI/DI 10195327226589 (each), 40195327226580 (case), Lot	US Nationwide distribution in the states of IN, KY, CA, MI, TX, WA. 111 distributed in the State of California.	May 2025 and prior.

	Number: 24LDA268; 6) REF ACC010898: UDI/DI 10198459192074 (each), 40198459192075 (case), Lot Number: 24LDA260.		

For additional information, please visit the FDA Website.

Confidential - Low