California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

B Braun Medical Inc. Recalls B.Braun Spinocan Spinal Needle

Recall Date	Product Description	Recalling Firm	Recall Reason
6/3/2026	Spinocan Spinal Needle procedure kits: Material Description (Material Number): Spinocan 26 Ga. x 3-1/2 in. (90 mm), Bupivacaine 0.75% with Dextrose 8.25% Tray (Kit) (333201); Spinocan 25 Ga. x 3-1/2 in. (90 mm), Bupivacaine 0.75% with Dextrose 8.25% Tray (Kit) (333211); Spinocan 22 Ga. x 3-1/2 in. (90 mm), Bupivacaine 0.75% with Dextrose 8.25% Tray (Kit) (333231)	B Braun Medical Inc. Bethlehem, PA	These procedure kits contain Bupivacaine Hydrochloride in Dextrose Injection, USP. The manufacturer of the Bupivacaine, Huons Co, Ltd., has issued a voluntary recall due to potential quality issues identified during a recent FDA manufacturing facility inspection.

Recall Class	Product Identification	Distribution	Affected Dates
I	Material: 333201; UDI-DI Primary/Unit of Use: 4046964179365/4046964179358; Batch Number: 0061976152;	16,080 Units Nationwide	May 2026 and prior

	Material: 333211; UDI-DI Primary/Unit of Use: 4046964179389/4046964179372; Batch numbers: 0061965729, 0061971323, 0061972572; Material: 333231; UDI-DI Primary/Unit of Use: 4046964179488/4046964179471; Batch Number: 0061978177		

For additional information, please visit the FDA website