State of California—Health and Human Services Agency
California Department of Public Health

Food and Drug Branch – Drug Recalls

California Drug Recall Information Sheet

FDA warns consumers

Baxter Issues Voluntary Nationwide Recall of One Lot of Heparin Sodium 0.9% Sodium Chloride Injection Due to Potential for Elevated Endotoxin Levels

Recall Date	Product Description	Recalling Firm	Recall Reason
08/06/2024	Heparin Sodium in 0.9% Sodium Chloride Injection	Baxter International Inc. (NYSE:BAX)	Use of heparin with higher than acceptable endotoxin levels may lead to significant adverse health consequences ranging from febrile reactions to toxic shock, multi-organ failure and death.

Recall Class	Product Identification	Distribution	Affected Dates
N/A	**Packaging:** 2,000 USP units, 1,000 mL in VIAFLEX Plus Plastic Container-1 unit per pouch **Lot number**: N008235 **NDC Number**: 0338-0433-04	Nationwide	Distributed between March 12, 2023, and August 24, 2023

For additional information, please visit the FDA website.

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