



California Drug Recall Information Sheet

FDA warns consumers

B. Braun Issues Voluntary Nationwide Recall of 0.9% Sodium Chloride for Injection USP 1000 mL in E3 Containers Due to the Potential for Particulate Matter and Leakage

Recall Date	Product Description	Recalling Firm	Recall Reason
08/08/2024	0.9% Sodium Chloride for Injection USP 1000 mL in E3 containers	B. Braun Medical Inc.	There is a potential for particulate matter and fluid leakage of the containers. This carries a risk for a reasonable probability of embolic phenomena such as stroke or ischemia/infarct to other organs and possible infection if these particulates are not sterile that could lead to permanent damage or impairment of body function which could be life-threatening.

Recall Class	Product Identification	Distribution	Affected Dates
N/A	Packaging: NACL INJ 0.9% 1000ML – E8000 Expiration date: 31 March, 2025 Lot number: J2L763, J2L764 NDC Number: 0264-7800-09	Nationwide	Distributed between 01 February 2024 – 28 February 2024

For additional information, please visit the [FDA website](#).

