California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Draeger, Inc. – Vapor 2000 Anesthetic Vaporizer Recall

Recall Date	Product Description	Recalling Firm	Recall Reason
12/23/2025	Draeger Vapor 2000, an unheated, calibrated anesthetic vaporizer used to deliver anesthetic agents during anesthesia administration.	Draeger, Inc. Telford, PA, USA	A certain component in affected Vapor 2000 anesthetic vaporizers was not delivered within specification and contained impurities, which could affect device safety and performance.

Recall Class	Product Identification	Distribution	Affected Dates
I	**Lot Code:** N/A; **Model No.:** M35054 **UDI-DI:** 04048675228059 **S/N range:** ASUK-0421 through ASUK-1532 (ranges)	431 Units; Nationwide and Worldwide Distribution including California	November 2025 and prior.

For additional information, please visit the FDA Website.