



Food and Drug Branch – Device Recalls

**CALIFORNIA DEVICE RECALL INFORMATION SHEET**

**Draeger Medical Recalls Carina Subacute Care Ventilator For Presence Of PE-PUR And Emission Of Dichloropropanol**

Recall Date	Product Description	Recalling Firm	Recall Reason
8/30/2023	<b>Carina Sub-Acute Care Ventilator</b>	<b>DRAEGER MEDICAL, INC.</b> Telford, Pennsylvania	Presence of poly-ether polyurethane (PE-PUR) and emission of 1,3-Dichloropropan-2-ol that exceed the acceptable uptake level during continuous use (>30 days) in pediatric patients.

Recall Class	Product Identification	Distribution	Affected Dates
I	<b>Carina Sub-Acute Care Ventilator</b> Part No. 5704110; UDI-DI 04048675398516; All Serial No.	11621 Units Nationwide	July 2023 and prior

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE [FDA WEBSITE](#)

