



Food and Drug Branch – Device Recalls

**CALIFORNIA DEVICE RECALL INFORMATION SHEET**

**Drager Medical Recalls SafeStar 55 for Possible Occlusion of Filters**

Recall Date	Product Description	Recalling Firm	Recall Reason
5/16/2022	<b>Drager SafeStar 55</b> Catalog No. MP01790	<b>Draeger Medical, Inc.</b> Telford, Pennsylvania	Possible occlusion of filters due to manufacturing error.

Recall Class	Product Identification	Distribution	Affected Dates
I	UDI-DI: 04048675026785 Lot No. LT2103.	429 Batteries Nationwide including California	May 2022 and prior

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

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