California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

The Metrix Company Imed Products Imed EVA BAG

Recall Date	Product Description	Recalling Firm	Recall Reason
11/8/2024	Imed Products Imed EVA BAG, 500 mL, 2-PORTS, REF IM68050; An empty single-use pouch made of plastic intended to contain a fluid for intravenous (IV) administration.	The Metrix Company	A limited number of IV bags have been found to leak during filling.

Recall Class	Product Identification	Distribution	Affected Dates
II	UDI/DI 00812496011329, Lot Numbers: 68050-A8320, 68050-A8435, 68050-A8476	864 cases of 50 units Worldwide - US Nationwide Including CA	November and Prior

For additional information, please visit the FDA Website.