California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

C.R. Bard Inc Lumen Esophagogastric Tamponade, Esophageal-Nasogastric, and Intragastric Linton Balloon Tubes

Recall Date	Product Description	Recalling Firm	Recall Reason
05/20/2025	Bard Minnesota Four Lumen Esophagogastric Tamponade Tube REF: 0092220 Bard Blakemore Esophageal-Nasogastric Tube (Adult) REF: 0092100 Bard Blakemore Esophageal-Nasogastric Tube (Child) REF: 0092110 Bard Blakemore Esophageal-Nasogastric Tube (Intermediate) REF: 0092300 Bard Single Intragastric Linton Balloon Tube REF: 0092740	C.R. Bard Inc	Due to customer complaints, there is a potential for users to be unable to or find it difficult to remove the plastic plugs in order to inflate the gastric and/or esophageal balloons.

Recall Class	Product Identification	Distribution	Affected Dates
I	All Lots within expiration UDI: (01)00801741076824 All Lots within expiration	N/A	April and prior

	UDI: (01)00801741076800 All Lots within expiration UDI: (01)00801741076817 All Lots within expiration UDI: (01)00801741076831 All Lots within expiration UDI: (01)00801741076848		

For additional information, please visit: FDA Website (Lumen Esophagogastric Tamponade), FDA Website (Esophageal-Nasogastric Adult), FDA Website (Esophageal-Nasogastric Child), FDA Website (Esophageal-Nasogastric Intermediate), and FDA Website (Intragastric Linton).