State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

CareFusion 303 Recalls Bd Alaris PCU for Containing Syringes that Have not Been Validated

Recall Date	Product Description	Recalling Firm	Recall Reason
11/15/2023	**Bd Alaris Pcu** REF 8015	**CAREFUSION 303, INC.** San Diego, California	Alaris PCA Module 8120, Patient Controlled Analgesia infusion pump: compatible syringes labeling contains syringes that have not been validated.
11/15/2023	**Bd Alaris Syringe Module** REF 8110	**CAREFUSION 303, INC.** San Diego, California	Alaris PCA Module 8120, Patient Controlled Analgesia infusion pump: compatible syringes labeling contains syringes that have not been validated.
11/15/2023	**Alaris Pca Module 8120**	**CAREFUSION 303, INC.** San Diego, California	Alaris PCA Module 8120, Patient Controlled Analgesia infusion pump: compatible syringes labeling contains syringes that have not been validated.

Recall Class	Product Identification	Distribution	Affected Dates
I	**BD Alaris PCU** All Lots/ UDI DI: 10885403516030,10885403515316,10885403812033,10885403515286,10885403812026,	867,362 Units Nationwide including California	15 September 2023 and prior

	10885403515293,10885403494291,10885403515309,10885403812002		
I	**BD Alaris Syringe Module** All Lots/ UDI-DI:10885403516047, 10885403515323, 10885403811043, 10885403515255 10885403811036, 10885403515262, 10885403811012, 10885403515279, 10885403424267	133,727 Units Nationwide including California	15 September 2023 and prior
I	**Alaris Pca Module 8120** All Lots/ UDI-DI: 10885403516023,10885403515231,10885403801549,10885403515248, 10885403801532,10885403515224,10885403801518	86,393 Units Nationwide including California	15 September 2023 and prior

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