State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

Becton Dickinson (BD)/Carefusion 303 Recalls Alaris Infusion Pumps Due To Compatibility Issues With Cardinal Health Monoject Syringes

Recall Date	Product Description	Recalling Firm	Recall Reason
12/1/2023	**Bd Alaris Syringe Module** REF 8110	**CAREFUSION 303, INC.** San Diego, California	Incompatible syringe sizes and models with the BD Alaris Syringe and PCA Modules can impact syringe pump operation resulting in inaccurate fluid delivery, delayed generation of occlusion alarms, and other potential problems
12/1/2023	**Bd Alaris Pcu** REF 8015	**CAREFUSION 303, INC.** San Diego, California	Incompatible syringe sizes
12/1/2023	**Alaris Pca Module** REF 8120	**CAREFUSION 303, INC.** San Diego, California	Incompatible syringe sizes

Recall Class	Product Identification	Distribution	Affected Dates
I	**Bd Alaris Syringe Module** All Lots/ UDI-DI: 10885403516047, 10885403515323, 10885403811043, 10885403515255 10885403811036, 10885403515262,	133727 Units Nationwide including California	September, 2023 and prior.

	10885403811012, 10885403515279, 10885403424267		
I	**Bd Alaris Pcu** All Lots/ UDI DI: 10885403516030, 10885403515316, 10885403812033, 10885403515286, 10885403812026, 10885403515293, 10885403494291, 10885403515309, 10885403812002	867362 Units Nationwide including California	September, 2023 and prior.
I	**Alaris Pca Module** All Lots/ UDI-DI: 10885403516023, 10885403515231, 10885403801549, 10885403515248, 10885403801532, 10885403515224, 10885403801518	86,393 Units Nationwide including California	September, 2023 and prior.

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE or BD Support Center at: 888-562-6018.