California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

CareFusion 303, Inc. BD Alaris Systems Manager and BD Care Coordination Engine Infusion Adapter

Recall Date	Product Description	Recalling Firm	Recall Reason
03/13/2025	BD Alaris Systems Manager, REF 9601, Infusion Safety Management Software BD Care Coordination Engine (CCE) Infusion Adapter, Safety Management Software	CareFusion 303, Inc.	Software issue that may result in outdated automated programming request(APR) being sent to the progressive care unit (PCU).

Recall Class	Product Identification	Distribution	Affected Dates
I	Software v12.5.1 or v12.5.2, and prior versions (4.33, 12.0.1, 12.0.2, 12.1, 12.1.2, 12.3) UDIs: 10885403960123 10885403519666 (10885403960116) Software: v1.7.2 UDI: 10885403510472	N/A	February and prior

For additional information, please visit: FDA Website (Alaris) and FDA Website (Care).