



California Medical Device Recall Information



Recall Name

CareFusion 303 Recalls Alaris Pump Module (Model 8100) Due to Potential Software Failure

Recall Date	Product Description	Recalling Firm	Recall Reason
04/23/14	Alaris Pump Module (Model 8100) Software Version 9.1.18	CareFusion 303, Inc. San Diego, CA	<i>A software failure may cause the pump module to not properly delay an infusion when the "Delay Until" option or "Multidose" feature is used.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	Affected Serial Numbers	CA, nationwide	Manufactured from: February 6, 2014 to April 8, 2014 Distributed from: February 7, 2014 to April 7, 2014.

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm398178.htm>