



# California Medical Device Recall Information



## Recall Name

### CareFusion Recalls Alaris Syringe Pump Due to Potential Malfunction of the Pump Alarm Error

Recall Date	Product Description	Recalling Firm	Recall Reason
7/02/15	Alaris Syringe Pump <ul style="list-style-type: none"><li>• <b>Model No. 8110</b></li></ul>	<b>CareFusion 303, Inc.</b> San Diego, CA	<i>Potential malfunction in the user interface when the user clears the syringe pump error code 351.6740.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	<a href="#">Affected Serial Numbers</a>	<b>CA</b> , nationwide	Manufactured and distributed from:  March 17, 2014 to September 30, 2014

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm460102.htm>