



# California Medical Device Recall Information



## Recall Name

### Hospira Recalls the GemStar Infusion System Due to Potential Battery Failure

Recall Date	Product Description	Recalling Firm	Recall Reason
3/18/13	GemStar Infusion System,  Models: 13000, 13100, 13150, 13086, 13087, 13088	<b>Hospira, Inc.</b> Lake Forest, IL	<i>Potential for battery failure to cause delay/interruption in therapy.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	GemStar Infusion System  Models Recalled: <ul style="list-style-type: none"> <li>13000, 13100, 13150, 13086, 13087, 13088</li> </ul> All serial numbers recalled	<b>CA</b> , nationwide	Manufacture and Distribution Dates: February 1999 through April 2013

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm349866.htm>