



# California Medical Device Recall Information



## Recall Name

### Hospira Recalls the Symbiq Infusion System Due to Potential Touchscreen Malfunction

Recall Date	Product Description	Recalling Firm	Recall Reason
8/29/12	Symbiq Infusion System, Models 16026 and 16027	<b>Hospira, Inc.</b> Lake Forest, IL	<i>Potential for Touchscreen Malfunction</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	Symbiq Infusion System  Models Recalled: <ul style="list-style-type: none"> <li>• 16026 Symbiq One Channel Infuser</li> <li>• 16027 Symbiq Two Channel Infuser</li> </ul> All serial numbers recalled	<b>CA</b> , nationwide	All

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm326020.htm>