



California Drug Recall Information



Recall Name

**Hospira Recalls Sodium Bicarbonate Injection
Due to Presence of Particulate**

Recall Date	Product Description	Recalling Firm	Recall Reason
03/18/16	8.4% Sodium Bicarbonate Inj., USP NDC: 0409-6625-02	Hospira, Inc. Lake Forest, IL	<i>Due to the presence of a particulate within a single-dose glass fliptop vial.</i>
Recall Class	Product Identification	Distribution	Affected Dates
N/A	Lot 56-148-EV Expiry 1AUG2017	CA , nationwide	Expiry 1AUG2017

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/Recalls/ucm491476.htm>