



California Drug Recall Information



Recall Name

Hospira Recalls Bupivacaine HCL Injection Due to Presence of Particulate Matter

Recall Date	Product Description	Recalling Firm	Recall Reason
Initial: 7/12/13 Expanded: 8/29/13	<ul style="list-style-type: none"> 0.25% Bupivacaine HCL Injection, USP (2.5 mg/ml), 30 ml single-dose vial NDC 0409-1159-02 0.75% Bupivacaine HCL Injection, USP (7.5 mg/ml), 30 ml single-dose vial NDC 0409-1165-02 	Hospira, Inc. Lake Forest, IL	<i>Confirmed customer reports of particulate floating and/or embedded in the glass vial.</i>
Recall Class	Product Identification	Distribution	Affected Dates
N/A	<p>0.25% Bupivacaine HCL Injection, USP (2.5 mg/ml):</p> <p>Lot# 18-136-DK* Exp. Date 1JUN2014</p> <p>0.75% Bupivacaine HCL Injection, USP (7.5 mg/ml):</p> <p>Lot# 23-338-DK* Exp. Date 1NOV2014</p> <p>Both packaged in 25 units per carton / 50 units per case, in glass tear-top vials.</p>	CA , nationwide	<p>Lot 18-136-DK distributed: August 2012 through September 2012</p> <p>Lot 23-338-DK distributed: January 2013 through May 2013</p>

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/Recalls/ucm368401.htm>