



# California Drug Recall Information



## Recall Name

**Hospira Recalls 1% Lidocaine HCL Injection  
Due to Presence of Dark Particulate**

Recall Date	Product Description	Recalling Firm	Recall Reason
10/04/13	<b>1% Lidocaine HCL Injection USP, 10 mg/ml, 20mL Vial</b>  <b>NDC 0409-4276-01</b>  Multiple-dose Flip-top Vial	<b>Hospira, Inc.</b> Lake Forest, IL	<i>Confirmed reports of visible particulates.</i>
Recall Class	Product Identification	Distribution	Affected Dates
N/A	Lot recalled: <ul style="list-style-type: none"><li>• <b>Lot #25-090-DK</b> (this number may be followed by "01" or "02")</li></ul>	<b>CA</b> , nationwide	Distributed between:  March 2013 and June 2013

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/Recalls/ucm370780.htm>