



# California Drug Recall Information



## Recall Name

### Hospira Recalls Lidocaine HCl Injection Due to Particulate Matter

Recall Date	Product Description	Recalling Firm	Recall Reason
07/29/14	Lidocaine HCl Injection, USP, 2% 20 mg per mL Single-dose Vial, Preservative-free  <b>NDC # 0409-2066-05</b>	<b>Hospira, Inc.</b> Lake Forest, IL	<i>Due to a confirmed customer report of discolored product with visible particles in the solution and embedded in the molded glass container.</i>
Recall Class	Product Identification	Distribution	Affected Dates
N/A	<b>Lot 25-550-DD</b>  <b>Expires 1JAN2015</b>	<b>CA</b> , nationwide	Distributed from:  June 2013 through July 2013.

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/Recalls/ucm407169.htm>