



California Drug Recall Information



Recall Name

Hospira Recalls Heparin Sodium Injection Due to Particulate Matter

Recall Date	Product Description	Recalling Firm	Recall Reason
9/11/14	Heparin Sodium, 1,000 USP Heparin Units, in 0.9% Sodium Chloride Injection, 500mL NDC # 0409-7620-03	Hospira, Inc. Lake Forest, IL	<i>Due to a confirmed customer report of particulate in a single unit. The foreign particulate was confirmed as human hair.</i>
Recall Class	Product Identification	Distribution	Affected Dates
N/A	Lot 41-046-JT Expires 01NOV2015	CA, nationwide	Distributed from: June 2014 through August 2014.

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/Recalls/ucm414201.htm>