



California Drug Recall Information



Recall Name

Hospira Recalls Metoclopramide and Ondansetron Injections Due to Presence of Glass Particulate Matter

Recall Date	Product Description	Recalling Firm	Recall Reason
10/02/13	<p>1. Metoclopramide Injection, USP (5 mg/ml) 2 ml single-dose vial NDC 0409-3414-01</p> <p>2. Ondansetron Injection, USP (2 mg/ml) 2 ml single-dose vial NDC 0409-4755-03</p> <p>Both products are packaged in 25 units per carton / 100 units per case, in glass flip-top vials.</p>	Hospira, Inc. Lake Forest, IL	<i>Confirmed vial defect where glass particulate matter (glass strands) were identified as being affixed to the inside of the vial walls.</i>
Recall Class	Product Identification	Distribution	Affected Dates
N/A	<p>Metoclopramide Injection, USP (5 mg/ml):</p> <ul style="list-style-type: none"> Lot# 28-104-DK; Exp. October 1, 2014 <p>Ondansetron Injection, USP (2 mg/ml):</p> <ul style="list-style-type: none"> Lot# 29-484-DK Exp. May 1, 2015 Lot# 29-510-DK Exp. May 1, 2015 	CA, nationwide	<p>Distributed between:</p> <p>June 2013 and September 2013</p>

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm370663.htm>