California Drug Recall Information

Recall Name
Hospira Recalls Propofol Injectable Emulsion Due to Possible Particulate Matter

Recall Date	Product Description	Recalling Firm	Recall Reason
4/17/14	Propofol Injectable Emulsion, USP, 1% 200 mg/20 ml (10mg/ml), single-dose vial **NDC # 0409-4699-30**	**Hospira, Inc.** Lake Forest, IL	*Due to possible free-floating metal particulates.*

Recall Class	Product Identification		Distribution	Affected Dates
N/A	**Lot Number**	**Expiration Date**	**CA**, nationwide	Distributed between: August 2013 through December 2013
	29-614-DJ	1MAY2015		
	29-615-DJ	1MAY2015		
	29-616-DJ	1MAY2015		
	29-617-DJ	1MAY2015		
	29-628-DJ	1MAY2015		
	29-629-DJ	1MAY2015		
	29-630-DJ	1MAY2015		

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm393849.htm