California Drug Recall Information

Recall Name
Bristol-Meyers Squibb Recalls Coumadin (Warfarin Sodium) for Injection Due to Presence of Particulate Matter

Recall Date	Product Description	Recalling Firm	Recall Reason
6/30/14	Coumadin (Warfarin Sodium) for Injection 5mg Single Use Vials **NDC #: 0590-0324-35**	**Bristol-Meyers Squibb Company** New York, NY	*Possible presence of visible particulate matter.*

Recall Class	Product Identification	Distribution	Affected Dates
N/A		**CA**, nationwide	Distributed from: November 2011 to January 2014

Lot #	Expiration
00201125	Sep-2014
00201126	Nov-2014
00201127	Dec-2014
00201228	Jun-2015
00201229	Jul-2015
00201230	Sep-2015

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm403501.htm