



California Drug Recall Information



Recall Name

**Teva Pharmaceuticals Recalls Seven Lots of Amikacin Sulfate Injection USP,
500 mg/2mL (250 mg/mL) and 1 grm/4mL (250 mg/mL) Vials
Due to Glass Particulate**

Recall Date	Product Description	Recalling Firm	Recall Reason
08/02/16	Amikacin Sulfate Injection USP <ul style="list-style-type: none"> 500 mg/2mL (250 mg/mL) Vials 1 gram/4mL (250 mg/mL) Vials 	Teva Pharmaceuticals North Wales, PA	<i>Due to the potential for presence of glass particulate matter.</i>
Recall Class	Product Identification	Distribution	Affected Dates
N/A	Recalled Lot Numbers: <ul style="list-style-type: none"> 7080315, Exp. 3/2017 7400315, Exp. 3/2017 7410315, Exp. 3/2017 7980415, Exp. 4/2017 2381114, Exp. 11/2016 2771114, Exp. 11/2016 4760915, Exp. 9/2017 Click for Product NDC Numbers	CA, nationwide	Shipped between: July 10, 2015 and January 26, 2016

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/Recalls/ucm514656.htm>