California Drug Recall Information

Recall Name
Fresenius Kabi USA Recalls Benztropine Mesylate Injection, USP 2 mg/2 mL (1 mg/mL) Due to Glass Particles in the Vials

Recall Date	Product Description	Recalling Firm	Recall Reason
06/30/13	Benztropine Mesylate Injection, USP 2 mg/2 mL (1 mg/mL) [in 2 mL single dose vials]	**Fresenius Kabi USA** Lake Zurich, IL (distributor)	*Potential presence of glass particles in the vials.* *Administration of glass particulate poses life - threatening safety risks.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
N/A	APP Pharmaceuticals Label **NDC #63323-970-02:** • Lot #111412, expiry 11/2014 Nexus Pharmaceuticals Label **NDC #14789-300-02:** • Lot# 030712, expiry 03/2014 • Lot# 071212, expiry 07/2014 • Lot# 090512, expiry 09/2014 Product Labels	**CA**, nationwide	APP labeled product distributed: 02/06/13 to 05/31/13 Nexus labeled product distributed: 05/18/12 to 11/20/12

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm359299.htm