
From: Board of Pharmacy <pharmacy.subscriberlist@DCA.CA.GOV>
Sent: Monday, March 20, 2017 11:38 AM
To: PHARM-GENERAL@DCALISTS.CA.GOV
Cc: Pharmacy_Subscriberlist@DCA
Subject: Product Recall - Fresenius Kabi USA, LLC

Fresenius Kabi USA, LLC ("Fresenius Kabi"), formerly APP Pharmaceuticals, LLC, is recalling the following batches of Fluphenazine Decanoate Injection, USP 25 mg / mL, 5mL. fill in a 5mL. vial. This recall is due to out-of-specification results for Assay at the 13 month stability test station and limited to the following 4 product batches.

| Product Name/Size | NDC Number | Product Code | Batch Number | Expiration Date | First Ship Date | Last Ship Date |
|----------------------------------------------------------------------------|-------------------|---------------------|---------------------|------------------------|------------------------|-----------------------|
| Fluphenazine Decanoate Injection, USP 25 mg / mL, 5mL fill in a 5 mL vial. | 63323-272-05 | 27205 | 6111141 | 07/17 | 9/1/2015 | 12/10/2015 |
| | | | 6111222 | 08/17 | 12/10/2015 | 3/19/2016 |
| | | | 6112346 | 01/18 | 2/29/2016 | 4/26/2016 |
| | | | 6112725 | 03/18 | 8/16/2016 | 1/16/2017 |

Please immediately check your inventory, quarantine, discontinue distribution of the affected product, and contact your wholesaler for directions.