Ketorolac Tromethamine Injection, USP, 30 mg/mL, and Ketorolac Tromethamine Injection, USP, 60 mg/2 mL

Date: 4/20/2020

At Magellan Rx Management, we want to help you get the best possible care. We have created a site to share drug recall information.

Fresenius Kabi USA, LLC has posted a lot recall of Ketorolac Tromethamine Injection, USP, 30 mg/mL, and Ketorolac Tromethamine Injection, USP, 60 mg/2 mL.

About this Recall:
Fresenius Kabi USA, LLC is voluntarily recalling 13 lots of Ketorolac Tromethamine Injection, USP, 30 mg/mL, 1 mL fill in a 2 mL amber vial and Ketorolac Tromethamine Injection, USP, 60 mg/2 mL, 2 mL fill in a 2 mL amber vial to the user level due to the presence of particulate matter. These particles were found in 8 sample vials.

What this means to you:
Customers with questions regarding this recall may contact Fresenius Kabi at 1-866-716-2459 Monday through Friday, during the hours of 8:00 a.m. to 5:00 p.m. Central Time. The FDA is directing consumers to contact their physician or health care provider if they have experienced any problems that may be related to taking or using this drug product

For more information regarding this FDA Recall Notification, please refer to the FDA website:

https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/fresenius-kabi-issues-voluntary-nationwide-recall-13-lots-ketorolac-tromethamine-injection-usp-due

FDA contact information for reporting adverse events/quality complaints can be reached online at https://www.accessdata.fda.gov/scripts/medwatch/index.cfm?action=reporting.home

or by calling the FDA at 1-888-INFO-FDA (1-888-463-6332) and then select prompt #2

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