Ranitidine Products

Date: 4/1/2020

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

The FDA has posted a full market withdrawal of ALL Ranitidine Products.

About this Withdrawal:

The U.S. Food and Drug Administration (FDA) today announced it is requesting manufacturers withdraw all prescription and over-the-counter (OTC) ranitidine drugs from the market immediately. This is the latest step in an ongoing investigation of a contaminant known as N-Nitrosodimethylamine (NDMA) in ranitidine medications (commonly known by the brand name Zantac).

What this means to you:

Patients and consumers should contact their physician or health care provider with any questions related to this withdrawal and/or if they have experienced any problems that may be related to taking Ranitidine Products. The FDA recommends consumers stop taking OTC ranitidine drugs and consider other approved OTC products if they wish to continue treating their condition. The FDA also recommends if you have had a prescription for this drug, to talk with your doctor or pharmacist before stopping this drug.

Because of the current COVID-19 pandemic, the FDA recommends consumers not take their drugs to a drug take-back location but follow the specific disposal directions in the medication guide or package insert or follow the FDA's recommended steps, which include ways to safely dispose of these drugs at home.

For more information regarding this FDA Withdrawal Notification, please refer to the FDA website: https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts

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