

# Ranitidine Products

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

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

## About this Recall:

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 recall 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 Recall 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