

Drug recall notice for sucralfate tablets USP 1 gram

To assist you in the care of your patients, Humana is alerting you to the recall of sucralfate tablets USP 1 gram on July 11, 2025.¹ We recommend you review your medical records and contact all patients for whom you have prescribed this medication to warn them of the recall.

The drug manufacturer, Nostrum Laboratories, Inc., is voluntarily recalling these products due to the company filing for bankruptcy on Sept. 30, 2024. The company has ceased all operations. As a result of discontinuation of its Quality activities, Nostrum Laboratories cannot guarantee that any lots of this product still within expiry can meet all intended specifications through the product's labeled shelf life.

While specific risks to patients from use of this product cannot be identified or assessed with certainty, the company said it cannot rule out potential patient risks. No other Nostrum Laboratories products are impacted by this drug recall.

To date, Nostrum Laboratories has received no reports of adverse events related to this recall.

Medications included in this recall

Visit the [U.S. Food and Drug Administration \(FDA\) website](https://www.fda.gov) for specific details about the recalled medication.

Product name	National Drug Code	Lot number
sucralfate tablets USP 1 gram	29033-003-01	All lots within expiry
sucralfate tablets USP 1 gram	29033-003-05	All lots within expiry

Information for providers:¹

- We sent a letter to your Humana-covered patients with a claim for sucralfate tablets USP 1 gram and asked them to contact their providers if their medication is included in the recall and if they have experienced problems that could be related to using these drug products.
- Providers with questions can contact:
 - Nostrum Laboratories by email at recallcoordinator@nostrumlabsrecall.com.
- Patients can report adverse reactions or quality problems experienced with the use of these products to the FDA's MedWatch Adverse Event Reporting Program online, by phone or by fax.
 - **Online:** Submit the [report](#).
 - Select "Form FDA 3500 - Voluntary Reporting."
 - **Phone or fax:** Download the [form](#).
 - Complete and submit "Form FDA 3500 - Voluntary Reporting" by phone at 800-FDA-1088 (332-1088) or by fax to 800-FDA-0178 (332-0178).

Reference

1. "Nostrum Laboratories, Inc. Issues Voluntary Nationwide Recall of Sucralfate Tablets USP 1 Gram Within Expiry," U.S. Food and Drug Administration, last accessed July 14, 2025, <https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/nostrum-laboratories-inc-issues-voluntary-nationwide-recall-sucralfate-tablets-usp-1-gram-within>.