

Important Drug Recall Notice

TO ALL PARTICIPATING PHARMACIES

Circular Letter MC25-012-CG

June 10, 2025

FDA announced that **Amneal Pharmaceutical LLC**, is recalling three lots of Sulfamethoxazole/Trimethoprim Tablets, USP, 400 mg/80 mg to the consumer level as the tablets may exhibit black spots on the tablet surface due to microbial contamination. The observance of black spots was reported in a product quality complaint.

RECOMMENDATIONS

1. Amneal is notifying its customers by UPS and is arranging for return of all recalled products. Wholesalers/distributors are being asked to notify their customers of the recall and provide instructions to contact Amneal for the return of the recalled products to Amneal. Retailers are being asked to notify their customers and instruct consumers to contact Amneal directly for assistance with return of any recalled product and reimbursement information.
2. Individuals with questions regarding this recall can contact Amneal Pharmaceuticals by:
 - Phone: 833-582-0812 Monday-Friday, 8:00 am-5:00 pm, EST
 - Fax: 631-983-2595
 - E-mail to: sulfamethoxazole-trimethoprim-recall@amneal.com
3. For Medical Inquiries or to report Adverse Events, or quality problems experienced with the use of this product, please contact Amneal Drug Safety by phone at 1-877-835-5472, Monday - Friday, 8:00 am – 6:00 pm, EST, or e-mail at DrugSafety@amneal.com.
4. Consumers with questions regarding this recall can contact Stericycle at 1-888-275-0506, Monday – Friday, 9am – 5pm CST.
5. Consumers should contact their physician or healthcare provider if they have experienced any problems that may be related to taking or using this drug product.
6. Review your inventory to identify existence of recalled products.
7. Expect patients to visit your pharmacy to deliver recalled products and prepare your pharmacy staff on how to handle the situation.

MC-Rx Pharmacy Services Department

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Pharmacy Communications are available at: <https://apps.mc-rx.com/MCRx.Forms/Pharmacy.Communications/>

Amneal Pharmaceutical LLC Issues a Nationwide Recall of Sulfamethoxazole / Trimethoprim Tablets, USP, 400 mg/80 mg Only, Due to Microbial Contamination

SUMMARY:

Company Announcement Date:	June 02, 2025
FDA Publish Date:	June 04, 2025
Product Type:	Drugs
Reason for Announcement:	Microbial contamination
Company Name:	Amneal Pharmaceutical LLC
Brand Name:	Amneal
Product Description:	Sulfamethoxazole/Trimethoprim Tablets, USP, 400 mg/80 mg

Company Announcement

Amneal Pharmaceutical LLC, is recalling three lots of Sulfamethoxazole/Trimethoprim Tablets, USP, 400 mg/80 mg to the consumer level as the tablets may exhibit black spots on the tablet surface due to microbial contamination. The observance of black spots was reported in a product quality complaint.

Risk Statement: Oral products contaminated with *Aspergillus* may result in serious and life-threatening infections. The use of the defective product in patients with underlying immunosuppressive conditions increases the concern for serious infections. To date, Amneal Pharmaceuticals has received no reports of adverse events, illnesses or injuries related to this recall.

The recalled product was distributed nationwide to wholesalers/distributors between the dates of 12/4/2024 to 5/15/2025 only.

The product is indicated for the treatment of Urinary tract infections caused by susceptible strains of the following organisms: *Escherichia Coli*, *Klebsiella* species, *Enterobacter* species, *Morganella morganii*, *Proteus mirabilis* and *Proteus vulgaris*. Acute otitis media in pediatric patients. Acute exacerbations of chronic bronchitis due to susceptible strains of *Streptococcus*

pneumoniae. Enteritis caused by susceptible strains of *Shigella flexneri* and traveler's diarrhea in adults.

Sulfamethoxazole/Trimethoprim Tablets, USP, 400 mg/80 mg is packaged in 100 tablet count and 500 tablet count bottles. This recall pertains only to the **400 mg/80 mg strength** and only to the listed Lots below. The lot number can be found on the Amneal bottle label or consult your pharmacy if you received a pharmacy vial. No other Sulfamethoxazole/Trimethoprim Tablets, USP, 400 mg/80 mg lots are impacted.

Lot Number	NDC Number	Expiration date	Date of First Distribution	Bottle Pack Size
AM241019	65162-271-10	06/2027	12/4/2024	100 count
AM241019A	65162-271-50	06/2027	12/4/2024	500 count
AM241020	65162-271-10	06/2027	12/4/2024	100 count

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Adverse reactions or quality problems experienced with the use of this product may be reported to the FDA's MedWatch Adverse Event Reporting program either online, by regular mail or by fax.

- Complete and submit the report [Online](#)

- Regular Mail or Fax: Download form or call 1- 800-332-1088 to request a reporting form, then complete and return to the address on the pre-addressed form, or submit by fax to 1-800-FDA-0178