



Phone: (614) 492-8177
Toll Free Phone: (800) 707-4621

URGENT PRODUCT RECALL

October 6, 2026

Dear Valued Customer,

American Health Packaging, Inc. is initiating a voluntary drug recall to the **RETAIL LEVEL** for **AHP Metoclopramide Tablets 5 mg, 100 UD**; **Carton NDC#: 60687-620-01, (Individual Dose NDC: 60687-620-11)**, for the lot listed below:

Product Description		AHP Lot No.	Expiration Date	Ship Dates of Product
AHP Metoclopramide Tablets 5 mg, 100 UD Carton NDC#: 60687-620-01 (Individual Dose NDC: 60687-620-11)		1028135	07/31/2027	1/9/2026 to 3/9/2026
		1029555	07/31/2027	3/9/2026 to 4/27/2026
REASON	This recall is being initiated in support of the recall by the manufacturer (Teva Pharmaceuticals USA, Inc) dated September 29, 2026, which included lots that were repackaged by American Health Packaging. These lots are being recalled as a safety precaution because, to date, a single Buspirone Hydrochloride Tablet, USP (10 mg) was discovered in each of two individual sealed bottles of Metoclopramide Tablets USP, 5mg lot # 8636085.			
PRODUCT INDICATION	Metoclopramide tablets are indicated for the: Treatment for 4 to 12 weeks of symptomatic, documented gastroesophageal reflux in adults who fail to respond to conventional therapy; Relief of symptoms in adults with acute and recurrent diabetic gastroparesis.			
HEALTH HAZARD EVALUATION	Teva’s health hazard assessment concluded that the main potential consequences are omission of the intended dose or inadvertent ingestion of a foreign tablet. While accidental administration of a single buspirone tablet would not be expected to result in clinically significant consequences for most patients, serious adverse outcomes cannot be excluded in susceptible individuals, including patients with buspirone hypersensitivity, patients receiving interacting medications (e.g., MAO inhibitors or serotonergic agents), or patients with relevant hepatic or renal impairment. Therefore, considering the most susceptible patient population, the severity of the hazard was assessed as severe, the likelihood of the harm occurrence is remote. Consequently, the overall risk of harm in patient population is considered to be medium.			
ACTIONS REQUIRED				
1. Distributors/Pharmacies – Immediately examine your inventory, quarantine and discontinue distribution of this lot. 2. Distributors – <u>Complete the enclosed Business Reply Card even if you do not have any product on hand.</u> 3. Distributors – Please pass this Recall Notice on <u>ONLY</u> to pharmacies that received this product lot. 4. Pharmacies – If you have units of the affected products/lot in inventory, please contact Sedgwick at (866)-737-4762 to receive a notification package with the Business Reply Card and return instructions. 5. Business Reply Cards can be submitted by any of these methods. Fax: (866)-891-0186 Email: ahp6949@sedgwick.com Mail: 2670 Executive Dr., Ste. A, Indianapolis, IN 46241 6. Distributors/Pharmacies – Return recalled product to Sedgwick as instructed in recall/return packet. 7. Pharmacies - You do not need to contact any patients.				
OTHER	This Recall extends to the Retail Level only. No other lots, packages, or formulations are being recalled. Please reorder stock immediately. For questions about the recall process, call Sedgwick at (866)-737-4762. This recall is being conducted with the knowledge of the Food and Drug Administration. We appreciate your immediate attention and cooperation and sincerely regret any inconvenience caused by this action.			

To receive credit, the reply form and recalled product must be returned to Sedgwick by January 31, 2027.

Thank you for your support in complying with the requests in this letter. We apologize for the inconvenience that this incident may cause you or your customers.

Sincerely,

Becky Mahon (Oct 5, 2026 14:17:01 EDT)

Becky A Mahon

Sr. Regulatory Specialist

American Health Packaging



AMERICAN
HEALTH
PACKAGING®

Phone: (614) 492-8177
Toll Free Phone: (800) 707-4621

Carton Label

NDC 60687-**620**-01

Metoclopramide
Tablets, USP

5 mg

100 Tablets (10 x 10)

Rx Only



(01) 0 03 60687 620 01 5

PHARMACIST: Dispense with Medication Guide to each patient.

NDC 60687-**620**-01

Metoclopramide
Tablets, USP

5 mg

100 Tablets (10 x 10)

Rx Only

Each Tablet Contains:
Metoclopramide hydrochloride, USP equivalent to 5 mg metoclopramide.

Usual Dosage: See full prescribing information.

Store at 20° to 25°C (68° to 77°F); excursions permitted between 15° to 30°C (59° to 86°F) [see USP Controlled Room Temperature].

PROTECT FROM LIGHT.

FOR YOUR PROTECTION: Do not use if blister is torn or broken.

The drug product contained in this package is from NDC # 0093-2204, Teva Pharmaceuticals.

Distributed by: American Health Packaging, Columbus, Ohio 43217

Scan for current
Medication Guide and
Prescribing Information →



762001
0462001/0824

Blister Label

Metoclopramide
Tablet, USP
5 mg



(01) 003 60687 620 11 4
American Health Packaging, Columbus, Ohio 43217

Expiry: xx/xx
Lot: xxxxxx

Metoclopramide
Tablet, USP
5 mg



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Expiry: xx/xx
Lot: xxxxxx

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Tablet, USP
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(01) 003 60687 620 11 4
American Health Packaging, Columbus, Ohio 43217

Expiry: xx/xx
Lot: xxxxxx

2026-10-06 Metoclopramide Tablets Recall Letter (Teva) - Retail

Final Audit Report

2026-10-05

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"2026-10-06 Metoclopramide Tablets Recall Letter (Teva) - Retail" History

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