

The Board of Pharmacy has received notice of the following product recall. The Board strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.

American Health Packaging, Inc. is initiating a 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.

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.

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.

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.

Product Description	AHP Lot No,	Expiration Date	Ship Dates of Product
AHP Metoclopramide Tablets 5 mg, 100 UD	1028135	07/31/2027	1/9/2026 to 3/9/2026
Carton NDC#: 60687-620-01 (Individual Dose NDC: 60687- 620-11)	1029555	07/31/2027	3/9/2026 to 4/27/2026