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 voluntary drug recall to the RETAIL LEVEL for AHP Primidone Tablets USP, 50 mg, 100 UD; Carton NDC: 68084-202-01, (Individual Dose NDC: 68084-202-11), 250 mg, 100 UD; Carton NDC: 68084-203-01, (Individual Dose NDC: 68084-203-11).

This recall is being initiated in support of the recall by the manufacturer (Lannett Company, Inc) dated April,23, 2026, which included lots that were repackaged by American Health Packaging. This recall is being carried out due to a potential for cross-contamination with Acemetacin API due to an issue at the API manufacturer.

Primidone, used alone or concomitantly with other anticonvulsants, is indicated in the control of grand mal, psychomotor, and focal epileptic seizures. It may control grand mal seizures refractory to other anticonvulsant therapy.

A Health Hazard Evaluation was conducted to assess the risk of Primidone tablets contaminated with trace amounts of Acemetacin (an NSAID). Testing confirmed Acemetacin levels were below the limits of quantification, and the worst-case daily exposure (≤1.48 mg/day) is far below therapeutic doses (60-180 mg/day). The overall risk to patients is low, with no clinically relevant adverse effects expected.

Product Description	AHP Lot No.	Expiration Date	Ship Dates of Product
AHP Primidone Tablets USP 50 mg, 100 Tablets (10x10) Carton NDC: 68084-202-01 (Individual Dose NDC: 68084-202-11)	1028739 1025622	12/31/2027 06/30/2027	02/17/2026 to 03/24/2026 09/24/2025 to 11/10/2025
AHP Primidone Tablets USP 250 mg, 100 Tablets (10x10) Carton NDC: 68084-203-01 (Individual Dose NDC: 68084-203-11)	1027583	09/30/2027	01/14/2026 to 03/23/2026