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 is initiating a voluntary Market Withdrawal to the RETAIL LEVEL for AHP Oxycodone Hydrochloride Tablets (CII), 5 mg, 100 UD; Carton NDC#: 68084-354-01, (Individual Dose NDC: 68084-354-11), for the lots listed below.

American Health Packaging has received multiple customer complaints for card seal defects observed on the subject lots. Customers reported card seal defects (weak/non-existent seals), leading to the tablets falling out of their cavities.

Oxycodone hydrochloride is indicated for the management of pain severe enough to require an opioid analgesic and for which alternative treatments are inadequate.

Product Description	**AHP Lot No.**	**Expiration Date**	**Ship Dates of Product**
AHP Oxycodone Hydrochloride Tablets (CII) 5 mg, 100 UD Carton NDC#: 68084-354-01 (Individual Dose NDC: 68084-354-11)	1027932	06/30/2027	12/15/2025 to 12/18/2025
	1028360	08/31/2027	12/18/2025 to 01/05/2026