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 drug recall 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 lot listed below.

American Health Packaging has received multiple customer complaints for card seal defects observed on the subject lot. 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: AHP Oxycodone Hydrochloride Tablets (CII) 5 mg, 100 UD

NDC NUMBER: Carton NDC#: 68084-354-01 (Individual Dose NDC: 68084-354-11)

LOT NUMBER: 1032815

EXPIRATION DATE: 06/30/2028

SHIP DATES OF PRODUCT: 08/27/2026 to 10/20/2026