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 Oxycodone Hydrochloride Oral Solution USP, 5 mg/5 ml, CII, 40 LUD; Case NDC# 60687-406-77 (Individual Dose NDC 60687-406-40) and 50 LUD; Case NDC# 60687-406-67 (Individual Dose NDC 60687-406-40)** for the lots listed below:

Product Description	**AHP Lot No.**	**Expiration Date**	**Ship Dates of Product**
AHP Oxycodone Hydrochloride Oral Solution USP, 5 mg/5 mL, CII, 40 LUD **40 ct Case NDC# 60687-406-77**	1004276	11/30/2022	08/10/2021 to 09/21/2021
AHP Oxycodone Hydrochloride Oral Solution USP, 5 mg/5 mL, CII, 50 LUD **50 ct Case NDC# 60687-406-67**	1002771	11/30/2022	07/01/2021 to 02/23/2022
Individual Dose NDC# 60687-406-40	**10 ct Tray NDC# 60687-406-46**		
REASON	This recall is being initiated due to impurity failure at 0-time of the repackaged lot. Unknown impurity RRT 0.49 and Total Degradants results were above specification at the 0-time. This notification has been updated to include one (1) lot to this recall.		