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 Bupropion Hydrochloride Extended-Release Tablets USP (XL), 150 mg, 100 UD; Carton NDC#: 60687-782-01, (Individual Dose NDC: 60687-782-11). This recall is being initiated due to an out of specifications for dissolution at the initial stability time point. The product is dissolving faster than specified limits which could lead to a change in therapeutic efficacy. Bupropion hydrochloride extended-release tablets (XL) are indicated for the treatment of major depressive disorder (MDD), as defined by the Diagnostic and Statistical Manual (DSM). Bupropion hydrochloride extended-release tablets (XL) are indicated for the prevention of seasonal major depressive episodes in patients with a diagnosis of seasonal affective disorder (SAD).

Product Description	**AHP Lot No.**	**Expiration Date**	**Ship Dates of Product**
AHP Bupropion Hydrochloride Extended-Release Tablets USP (XL), 150 mg, 100 UD Carton NDC#: 60687-782-01 (Individual Dose NDC: 60687-782-11)	1017343	12/31/2025	03/15/2024 to 05/31/2024