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.

Hikma Pharmaceuticals USA Inc. (formerly West-Ward Pharmaceuticals) is initiating a drug recall of three (3) lots of **Buprenorphine Hydrochloride Injection, 0.3mg/mL, 1 mL vial** at the **Retail level.** This recall is being conducted with the knowledge of the Food and Drug Administration. Distribution dates: 04/28/2020 thru 05/26/2020.

Item Description	NDC	Lot	Expiration Date
Buprenorphine Hydrochloride Injection 0.3mg/mL (1 mL vial)	0143-9246-05	2005023.1	02/2021
Buprenorphine Hydrochloride Injection 0.3mg/mL (1 mL vial)	0143-9246-05	2005024.1	02/2021
Buprenorphine Hydrochloride Injectlon 0.3mg/mL (1 mL vial)	0143-9246-05	2005025.1	02/2021

Reason for Recall:

This recall is being conducted due to borderline and Out of Specification (OOS) assay results were found for Buprenorphine Hydrochloride Injection batches during the 3 months product stability testing.