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. and its affiliates ("Hikma") are initiating a drug recall of six lots of Ciprofloxacin Tablets, USP, 500 mg and 750 mg at the wholesale/distributor and retail pharmacy levels only.

This recall is being conducted due to N-nitroso-ciprofloxacin results above the FDA recommended interim control limit. A medical assessment has been completed and supports that the observed N-nitroso-ciprofloxacin levels are not expected to adversely impact patient safety. All other testing parameters for these lots are within specification.

This recall is limited to the six lot numbers listed below. No other "Hikma" labeled products, or lots, are impacted by this recall.

Product	**NDC No.**	**Lot No.**	**Exp. Date**	**Ship Dates to Customers**
Ciprofloxacin Tablets, USP, 500 mg 100 count bottle	0143-9928-01	CB496A	December 2026	09/11/2023
Ciprofloxacin Tablets, USP, 500 mg 100 count bottle	0143-9928-01	CB497A	December 2026	08/07/2023 - 09/11/2023
Ciprofloxacin Tablets, USP, 500 mg 100 count bottle	0143-9928-01	CB498A	December 2026	08/14/2023 - 01/08/2024
Ciprofloxacin Tablets, USP, 750 mg 50 count bottle	0143-9929-50	CA083A	December 2026	08/14/2023 - 05/16/2024
Ciprofloxacin Tablets, USP, 750 mg 50 count bottle	0143-9929-50	CA084A	December 2026	07/05/2023 - 11/10/2023

Ciprofloxacin Tablets, USP, 750 mg 50 count bottle	0143-9929-50	CA085A	December 2026	07/18/2023 - 03/27/2024