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.

Sun Pharmaceutical Industries, Inc. has initiated a recall in response to microbial contamination in stagnant water found in the duct of the manufacturing equipment. Shipment of this product initiated on September 8, 2023.

Product Name	Package Description	Lot Number	NDC Number	Expiration Date
Lurasidone Hydrochloride Tablets 60 mg	30 count	DNE0620A	47335-639-83	05/2025
Lurasidone Hydrochloride Tablets 120 mg	30 count	DNE0621A	47335-579-83	11/2024
Lurasidone Hydrochloride Tablets 120 mg	30 count	DNE0815A	47335-579-83	12/2024