The Board of Pharmacy has received notice of the following product recall:

Description: RANITIDINE HYDROCHLORIDE TABLETS, USP, 150 MG, 60's, 100's & 500's

Batch: 19190372
NDC#: 68462024860
Pack Size: 60's
Expiry Date: Dec-21
Date of Shipment: Jun-19

Batch: 19190372
NDC#: 68462024801
Pack Size: 100's
Expiry Date: Dec-21
Date of Shipment: Jun-19

Batch: 19190372
NDC#: 68462024805
Pack Size: 500's
Expiry Date: Dec-21
Date of Shipment: Jun-19

Recall has been initiated due to NDMA impurity found more than acceptable limits (0.32 ppm) for batch #19190372.

The Board of Pharmacy strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.