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

Description: AHP Ranitidine Tablets, USP, 150 mg 100 count Unit Dose Blisters Case NDC#: 60687-322-01 Individual Dose NOC: 60687-322-11 AHP Lot Number/Expiration Date/Ship Dates of Product:
179516 12/31/2019 07/30/2018 to 09/04/2018
179745 12/31/2019 09/04/2018 to 10/15/2018
180712 02/29/2020 10/11/2018to 11/12/2018
180819 04/30/2020 11/12/2018 to 12/11/2018
181403 05/31/2020 12/11/2018 to 01/31/2019
182544 05/31/2020 01/31/2019 to 05/31/2019
183155 05/31/2020 02/26/2019 to 04/11/2019
183236 05/31/2020 04/11/2019 to 05/13/2019
185739 12/31/2020 05/09/2019 to 08/02/2019
186600 12/31/2020 08/02/2019 to 09/12/2019
186702 12/31/2020 09/12/2019 to 10/02/2019

This recall is being initiated in support of the recall by the manufacturer (Amneal Pharmaceuticals, LLC) dated November 13, 2019, which included lots that were repackaged by American Health Packaging. Amneal stated that" This recall has been initiated due to the potential N-Nitrosodimethylamine (NDMA) amounts above levels established by the FDA. This recall is for all lots within expiration."

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.