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

Description: Ranitidine Oral Solution, USP 150 mg/10 mL unit dose product Lot #s/Exp Date:
501290/30-Nov-2019
501326/30-Nov-2019
501501/30-Nov-2019
501592/30-Apr-2020
501679/30-Apr-2020
NDC: 68094-204-59 / 68094-204-61 / 68094-204-62

Ranitidine Oral Solution, USP 150 mg/10 mL is being recalled due to the possible presence of N-nitrosodimethylamine (NDMA) impurity. This recall is being initiated in response to the recall by the manufacturer (Amneal Pharmaceuticals. LLC), which included affected lots that were repackaged by Precision Dose Inc.

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.