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

**LEVEL OF NOTIFICATION:** retail

**SUPPLIER:** American Health Packaging

| Description | Lot #/Exp Date | NDC | UPC |
|---|---|---|---|
| RANITID OS 150MG/10ML AHP UD40 | 187652 05/31/2020; 184278 10/31/2020; 183723 10/31/2020 | 60687026023 | 36068726023 |
| RANITID OS 150MG/10ML AHPUD50@ | 186563 03/31/2021; 184445 12/31/2020; 183449 10/31/2020; 178413 02/29/2020; 177874 01/31/2020 | 60687026069 | 36068726069 |

**NOTE: NDCs above for cases level of product. NDCs for individual product: for NDC 60687026023, Individual NDC: 60687026042. For NDC 60687026069, Individual NDC: 60687026042**

American Health Packaging is voluntarily recalling the above items due to the due to the presence of NDMA in the Ranitidine syrup. This recall is to the Retail level. Affected product started shipping October 16, 2018.

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.