

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

American Health Packaging is voluntarily recalling the above items 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.