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

Description	Lot # / Exp Date	NDC	UPC
RANITID SDV 25MG/ML 2ML ZYD10@	M812490 07/31/2020; M808367 04/30/2020; M808368 04/30/2020; M808376 04/30/2020; M809235 05/31/2020; M809238 05/31/2020; M812489 07/31/2020; M812491 07/31/2020; M812492 07/31/2020; M812933 07/31/2020; M813476 07/31/2020; M813770 07/31/2020; M906333 03/31/2020; M906334 03/31/2021; M809243 05/31/2020; M812742 07/31/2020	68382042202	36838242202
RANITID MDV 25MG/ML ZYD 6ML@	M818853 10/31/2020; M904264 02/28/2021; M904265 02/28/2021; M813773 07/31/2020	68382042306	36838242306
RANITIDINE INJ 25MG/ML ZYD 1	L900007 02/28/2021; L900005 02/28/2021; L900006 02/28/2021	70710155001	37071015501

Zydus is withdrawing the above items/lots based on FDA's recommendation due to impurity in some Ranitidine products increasing over time that may result in consumer exposure to unacceptable levels of the impurity. This withdrawal is to the retail level. Affected product started shipping November 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.