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

Description	Lot # / Exp Date	NDC	UPC
RANITIDINE CAP 150MG AJAN 60@	PA1849I 08/31/22; PA1859I 08/31/22; PA1879I 08/31/22; PA1319H 07/31/22; PA1469H 07/31/22; PA0519G 06/30/22; PA1139G 06/30/22; PA1119L 11/30/21; PA0659K 10/31/21; PA0679K 10/31/21; PA1209B 01/31/21; PA1859B 01/31/21; PA1098L 11/30/20; PA1128L 11/30/20; PA0968I 08/31/20; PA0988I 08/31/20; PA0998I 08/31/20; PA1008I 08/31/20; PA1018I 08/31/20	27241010906	32724110906
RANITIDINE CAP 150MG AJAN 500@	PA1869I 08/31/22; PA1879I 08/31/22; PA1329H 07/31/22; PA1339H 07/31/22; PA1349H 07/31/22; PA0519G 06/30/22; PA0529G 06/30/22; PA1139L 11/30/21; PA1149L 11/30/21; PA0639K 10/31/21; PA0649K 10/31/21; PA0659K 10/31/21; PA0669K 10/31/21; PA0679K 10/31/21; PA1209B 01/31/21; PA1859B 01/31/21; PA1339A 12/31/20; PA1108L 11/30/20; PA1118L 11/30/20; PA1128L 11/30/20; PA0968I 08/31/20; PA0978I 08/31/20; PA0998I 08/31/20; PA1018I 08/31/20	27241010950	32724110950
RANITIDINE CAP 300MG AJAN 30@	PA1559H 07/31/22; PA1569H 07/31/22; PA0539G 06/30/22; PA1519L 11/30/21; PA1529L 11/30/21; PA2009L 11/30/21; PA1849K 10/31/21; PA1539J 09/30/21; PA1549J 09/30/21; PA1219B 01/31/21; PA1229B 01/31/21; PA1028I 08/31/20; PA1038I 08/31/20; PA1048I 08/31/20	27241011003	32724111003
RANITIDINE CAP 300MG AJAN 100@	PA1569H 07/31/22; PA0539G 06/30/22; PA1529L 11/30/21; PA1549J 09/30/21; PA1219B 01/31/21; PA0768K 10/31/20; PA1048I 08/31/20	27241011010	32724111010

Ajanta is withdrawing the **all lots** of the above items as listed, due to laboratory test results suggesting that levels of NDMA in these products may increase above acceptable daily intake limits with temperature and time. This withdrawal is to the retail level. Affected product started shipping November 27, 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.