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

LEVEL OF NOTIFICATION: Retail

SUPPLIER: Major

Description	**Lot #s**	**NDC**	**UPC**
RANITID TAB 150MG MMP 50	8JE2154; 9AE2499; 9BE2863; 9CE3723; 9DE2891; 9EE2812; 8GE1921;	00904671651	30904671651
RANITID TAB 75MG MMP 30	9DE2721; 8JE1916; 8KE2243; 8ME2685; 9CE3317; 9EE2579; 9GE2785; 9AE2785	00904671546	30904671546
Ranitidine 150mg Tablets USP	**8GE1833; 9EE2760**	00904671624	**30904671624**
RANITID TAB 75MG MMP 60@	9GE2793; 8JE1917; 8KE2245; 8ME2724; 9AE2831; 9CE3339; 9DE2747; 9EE2636; 9FE2971	00904671552	30904671552

Major is recalling the above lots of Ranitidine due to possibility of presence of NDMA contaminants. This recall is to the retail level. Affected product started shipping August 9, 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.