

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

**LEVEL OF NOTIFICATION:** Retail pharmacy

**SUPPLIER:** Gericare

<b>Description</b>	<b>Lot #</b>	<b>NDC</b>	<b>UPC</b>
RANITIDINE TAB 75MG GERI 30@	All within expiry	57896076503	35789676503
Ranitidine Tablets USP 150mg	All within expiry	57896077024	
Ranitidine Tablets USP 150mg	All within expiry	57896077005	
Ranitidine Tablets USP 150mg	All within expiry	57896071724	
Ranitidine Tablets USP 150mg	All within expiry	57896071705	
Ranitidine Tablets USP 75mg	All within expiry	57896076506	
Ranitidine Tablets USP 75mg	All within expiry	57896071506	
Ranitidine Tablets USP 75mg	All within expiry	57896071503	

Gericare is recalling the above items due to the presence of NDMA levels found within the product in excess of amounts allowed by the FDA. This recall is to the retail pharmacy level.

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.