The Board of Pharmacy has received notice of the following product recall. The Board strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.

Description	Lot # and Exp Date	NDC	UPC
250MG/5ML	30210004 01-31-23; 30200015 05-31-22;	68462031732	36846231732
	30200016 05-31-22; 30200036 09-30-22;		
	30200038 09-30-22; 30200039 09-30-22;.		
	30200040 09-30-22; 30210001 12-31-22;		
	30200014 04-30-22; 30210003 01-31-23;		
	30210031 02-28-23; 30210005 01-31-23;		
	30210006 01-31-23; 30210014 02-28-23;		
	30210022 02-28-23; 30210028 02-28-23;		
	30210029 02-28-23; 30210030 02-28-23;		
	30210002 12-31-22		

Glenmark is recalling the above item/lots due to a lack of assurance of product sterility discovered after a review of the manufacturing facility's microbiology laboratory controls/processes. This recall is to the retail pharmacy level. Affected product started shipping September 2, 2020.