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.

Product: **CHLORZOXAZONE TABLETS USP 375MG 100'S (NDC #68462-724-01)** & **CHLORZOXAZONE TABLETS USP 750MG 100'S (NDC #68462-725-01)**

Batch	NDC	Pack size	Exp. Date
29200022	68462 -724-0l	375MG 100'S	3-31-2022
29200024	68462 -724-0 l	375MG 100'S	3-31-2022
29200035	68462 -724-0l	375MG 100'S	6-30-2022
29200023	68462-725-01	750MG 100'S	3-31-2022
29200025	68462-725-01	750MG 100'S	3-31-2022
29200036	68462-725-01	750MG 100'S	6-30-2022
29200056	68462-725-01	750MG 100'S	9-30-2022
29200070	68462-725-01	750MG 100'S	11-30-2022

Glenmark Pharmaceuticals Inc. USA is recalling to the **Retail Level** all unexpired lots **of CHLORZOXAZONE TABLETS USP, 375 mg (100 count) and 750 mg (100 count)**. This product is being recalled due to gaps in the quality system in the Quality Control microbiology laboratory. To date, Glenmark has not received any reports of adverse events related to this recall nor is there any confirmed contamination of the product.