

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: TELMISARTAN AND HYDROCHLOROTHIAZIDE TABLETS, USP, 80 MG / 12.5 MG, 3X10'S BLISTER PACK (NDC 68462-841-13) AND 80 MG / 25 MG, 3X10'S BLISTER PACK (NDC 68462-842-13)

Glenmark Pharmaceuticals is recalling at **Retail Level** the following batches listed in Table 1 below. Note: This recall is being extended from Wholesale to Retail level with the knowledge of FDA.

Table 1. Recall Batch Numbers

S. No.	NDC of Pack	NDC of Inner Blister	Batch No.	Expiry Date
1.	68462-842-13	6846284211	17210935	05/2023
2.	68462-842-13	6846284211	17210936	05/2023
3.	68462-842-13	6846284211	17211206	06/2023
4.	68462-842-13	6846284211	17211652	08/2023
5.	68462-842-13	6846284211	17211655	08/2023
6.	68462-842-13	6846284211	17211658	08/2023
7.	68462-841-13	6846284111	17210929	05/2023
8.	68462-841-13	6846284111	17210930	05/2023
9.	68462-841-13	6846284111	17211203	06/2023
10.	68462-841-13	6846284111	17211643	08/2023
11.	68462-841-13	6846284111	17211646	08/2023
12.	68462-841-13	6846284111	17211649	08/2023