

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.

This is to inform you of that Glenmark is initiating a recall to the Retail level involving Diltiazem Hydrochloride Extended Release Capsules 60 mg, 90 mg & 120 mg (100's Pack Container). The recall is to the retail level of the below identified product batches for Diltiazem Hydrochloride Extended Release Capsules has been initiated by Glenmark out of an abundance of precaution because the results of certain finished product analysis does not comply with the current recommended limit of "N-Nitroso-Desmethyl-Diltiazem" i.e. 0.074 ppm.

| S. No. | NDC | Batch # | Pack Size | Expiry |
|---------------|--------------|----------------|------------------|---------------|
| 1. | 68462-851-01 | 17222452 | 100 capsules | 11/2024 |
| 2. | 68462-562-01 | 17222470 | 100 capsules | 11/2024 |
| 3. | 68462-850-01 | 17222544 | 100 capsules | 11/2024 |
| 4. | 68462-562-01 | 17222547 | 100 capsules | 11/2024 |
| 5. | 68462-562-01 | 17230304 | 100 capsules | 12/2024 |
| 6. | 68462-562-01 | 17230598 | 100 capsules | 02/2025 |
| 7. | 68462-851-01 | 17230607 | 100 capsules | 02/2025 |
| 8. | 68462-562-01 | 17230680 | 100 capsules | 11/2024 |
| 9. | 68462-850-01 | 17230784 | 100 capsules | 03/2025 |
| 10. | 68462-850-01 | 17231080 | 100 capsules | 04/2025 |