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 notice is to inform you of a product recall involving **Diltiazem Hydrochloride Extended Release Capsules, 360 mg**

Product Name	Package Description	Lot Number	NDCNumber	Expiration Date
Diltiazem Hydrochloride Extended Release Capsules, 360mg	90 Count	HAC3120A	47335-679-8 1	04/2023
	90 Count	HAC3121A	47335-679-8 1	04/2023
	90 Count	HAC4460A	47335-679-8 1	10/2023
	90 Count	HAD0365A	47335-679-8 1	12/2023
	90 Count	HAD1452A	47335-679-8 1	02/2024

Sun Pharmaceuticals Inc. has initiated this recall due to related substance results (Deacetyl Diltiazem Hydrochloride Impurity) that were reported above the specification limit during stability testing; and not meeting to USP requirement for dissolution testing in FDA Laboratory. This recall should be carried out to the **retail** level.

Sun Pharmaceutical Industries Inc. initiated shipment of this product in November 2, 2021.