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.

Apotex Corp. is recalling two (2) batches of **Brimonidine Tartrate/Timolol Maleate Ophthalmic Solution** (batches VE0614 and VE0616) to the **Retail/Pharmacy level**. This recall is being initiated out of an abundance of caution. It follows the discovery of atypical weight loss in individual bottle from batch VE0614 that was stored horizontally. While this batch met all quality specifications at the time of release - and continues to meet all filed and approved stability test specifications - the company has decided to recall the aforementioned batches. The recall is also extended to include batch VE0616 as it was made with the same combination of components as batch VE0614.

Based on the completed health hazard assessment and test data conducted on the bottles with atypical weight loss, the observed atypical individual weight loss results will not have significant adverse effects on the safety and efficacy of the product and thus would not affect the risk benefit profile of drug product if used before expiry date. The details for the batches being recalled are listed below.

Product	Strength	Pack Size	NDC	UPC on Bottle	UPC on Carton	Batch Number	Expiry Date (mm/yyyy)	First Date of Sale (mm/dd/yyyy)	Last Date of Sale (mm/dd/yyyy)
Brimonidine Tartrate/Timolol Maleate Ophthalmic Solution	0.2% / 0.5%	5 mL	60505-0589-1	(01)00360505058914	360505058914	VE0614	12/2025	10/18/2024	05/16/2025
		10 mL	60505-0589-2	(01)0(03)60505058921	360505058921	VE0616	12/2025	07/31/2024	12/11/2024