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 VJ8599 and VL1668) to the **Retail/Pharmacy level.** Refer to Appendix A for the sample label of the product. This recall is being initiated out of an abundance of caution. It follows the discovery of atypical weight loss in individual bottles from batch VJ8599 that were 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 VL1668 as it was made with the same combination of component lots as batch VJ8599.

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 the drug product if used before expiry date. However, in a theoretical situation of sterility breach, patient safety may be affected. 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/Timlol Maleate Ophthalmic Solution	0.2%/ 0.5%	5mL	60505-0589-1	(01)00360505058914	36050558914	VJ8599	09/2026	04/10/2025	11/05/2025
						VL1668	01/2027	01/26/2026	02/20/2026