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.

Genentech is conducting a recall of CATHFLO® ACTIVASE®(alteplase) 2 mg vial, with knowledge of the U.S. Food and Drug Administration. Genentech is conducting this recall due to an ongoing investigation for deformed stoppers observed during filling operations for Cathflo Activase. A defective stopper may not seal properly resulting in a potential container closure integrity and sterility concern. To date, no safety issues have been reported. No other lots are affected.

Product Description	NDC	Lot Number	Expiration Date on box	Genentech Distribution Dates
CATHFLO® ACTIVASE® (alteplase) 2 mg vial		3618858	Jan 31, 2026	5/6/2024 - 5/9/2024
	50242-0041-64	3618873	Jan 31, 2026	4/26/2024 - 5/8/2024