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.

On July 7, 2022, Apotex Corp. notified impacted direct account Wholesalers and Distributors of a recall of two (2) lots of **Aripiprazole Tablets, USP 20mg** specified below. Apotex is expanding the Wholesaler/Distributor level recall for two (2) lots of **Aripiprazole Tablets, USP 20mg** specified below to a **Retail** level. Apotex is now notifying all impacted direct account Wholesalers, Distributors and Retailers of this recall up to the **Retail** level for these two (2) lots of Aripiprazole Tablets, USP 20mg. This recall is being conducted with the knowledge of the US FDA.

Product	Pack Size	Strength	NDC	UPC Code	Lot Number	Exp. Date (mm/yyyy)	First Date of Sale (mm/dd/ yyyy)	Last Date of Sale (mm/dd/yyyy)
Aripiprazole Tablets, USP	1000's Bottle	20mg	60505-2676-8	360505267682	TD9591	04/2023	08/30/2021	05/26/2022
	30's Bottle		60505-2676-3	360505267637	TD9592		07/23/2021	05/26/2022

REASON FOR MARKET ACTION: The recall is due to one out of specification dissolution result generated at the 12-month stability time point for one (1) tablet of lot TD9591. Out of an abundance of caution, lot TD9591 and lot TD9592 are included in the scope of this recall, since they are packaged from same bulk lot.