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.

Rising Health LLC has initiated a voluntary drug product recall for the product **Duloxetine DR Capsules USP 20 mg/30 mg/60 mg** from USA market ***due to presence of Nitrosamine Drug Substance Related Impurity (NDSRI), N-Nitroso-Duloxetine above the interim acceptable intake limit of 5ppm as stipulated by the FDA.***

This product was shipped between the dates of 03/16/2023 – 10/29/2024.

The recall involves the following batches of Duloxetine DR Capsules, USP 20mg, 30mg and 60mg:

NDC Number	Product Name and Dosage Strength	Package Size	Lot Number	Expiration Date
57237-017-60	Duloxetine DR Capsules USP 20mg	60's HDPE Bottle	DT2023003A	Jan-25
			DT2023007A	Jan-25
			DT2023008A	Jan-25
57237-018-99	Duloxetine DR Capsules USP 30mg	1000's HDPE Bottle	DT3023025A	Jan-25
57237-018-30	Duloxetine DR Capsules USP 30mg	30's HDPE Bottle	DT3023051A	Apr-25
57237-019-99	Duloxetine DR Capsules USP 60mg	1000's HDPE Bottle	DT6023002A	Dec-24
			DT6023048A	Jan-25
			DT6023016A	Dec-24
			DT6023036A	Dec-24
57237-019-30	Duloxetine DR Capsules USP 60mg	30's HDPE Bottle	DT6023053A	Jan-25
			DT6023061A	Jan-25
			DT6023068A	Jan-25
			DT6023074A	Jan-25
			DT6023078A	Feb-25
			DT6023076A	Feb-25
			DTC24043A	Dec-25
			DTC24044A	Dec-25