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.

This is to inform you that ***Breckenridge Pharmaceutical, Inc.*** (Breckenridge) is voluntarily performing a Retail Level Recall of **Duloxetine Delayed-Release Capsules, USP, 60mg,** manufactured by Towa Pharmaceutical Europe, S.L.

Only the lots listed in the table below are being recalled due to presence of Nitrosamine Drug Substance Related Impurity (NDSRI), N-nitroso-duloxetine, above the proposed interim limit.

Product	**Size**	**NOC Number**	**Affected Lot #**	**Exp Date**
Duloxetine Delayed-Release Capsules USP, 60 mg	1000-count	51991-748-10	240534C	01/2027
Duloxetine Delayed-Release Capsules USP, 60 mg	1000-count	51991-748-10	240977C	04/2027