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.

Please be advised that Breckenridge Pharmaceutical, Inc. (Breckenridge) is performing a Retail level Class II Recall of Duloxetine Delayed-Release Capsules, USP, 20mg, manufactured by Towa Pharmaceutical Europe, S.L. This is a Retail Level recall.

PRODUCT NAME: Duloxetine Delayed-Release Capsules USP, 20 me

NDC NUMBER: 51991-746-05

LOT NUMBER: 220128

EXPIRATION DATE: 12/2024