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.

Viatris Specialty LLC is conducting a recall at the retailer level of the below-listed lot of Xanax XR (alprazolam) Extended-release Tablets 3mg, which is packaged in a bottle containing 60 tablets.

This lot is being recalled out of an abundance of caution after testing some of the tablets produced out-of-specification dissolution results.

This lot was distributed in the United States between Aug 27, 2024, and May 29, 2025.

The risk to the patient associated with this event is considered to be negligible. To date, no reports of adverse reactions associated with this lot have been received.

Xanax XR is a benzodiazepine indicated for the treatment of panic disorder with or without agoraphobia, in adults.

This recall is being conducted with the knowledge of the Food and Drug Administration.

PRODUCT: Xanax XR (alprazolam) Extended-release Tablets 3mg, 60 Tablets in a Bottle

NDC NUMBER: 58151-506-91

LOT NUMBER: 8177156

EXPIRY DATE: Feb 2027