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 the following Lot No. 24291 of Nitroglycerin Injection, USP manufactured and distributed by American Regent, Inc. is the subject of a Recall. Recall of this product to RETAIL LEVEL was initiated due to particulate found during Reject Characterization.

Nitroglycerin Injection is indicated for treatment of peri-operative hypertension; for control of congestive heart failure in the setting of acute myocardial infarction; for treatment of angina pectoris in patients who have not responded to sublingual nitroglycerin and β-blockers; and for induction of intraoperative hypotension.

The administration of an injectable product containing particulate matter could potentially result in phlebitis, capillary occlusion, thromboembolism, or an immunologic or immunogenic response. When Nitroglycerin Injection, USP is used according to the approved prescribing procedures, at this time there is low potential risk of the particle entering the systemic circulation with consequent adverse reactions. To date, American Regent, Inc. has not received any reports of adverse events related to this recall.

PRODUCT: Nitroglycerin Injection, USP 50 mg/10 mL 10 mL/vial

NDC NUMBER: 0517-4810-25

LOT NUMBER: 24291

EXPIRATION DATE: August 31, 2026