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. 25090 of Selenious Acid Injection 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.

Selenious Acid Injection is indicated in adult and pediatric patients as a source of selenium for parenteral nutrition when oral or enteral nutrition is not possible, insufficient, or contraindicated.

The administration of an injectable product containing particulate matter could potentially result in phlebitis, capillary occlusion, thromboembolism, or an immunologic or immunogenic response. When Selenious Acid 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. As of July 24, 2026, American Regent, Inc. has not received any reports of adverse events related to this recall.

PRODUCT: Selenious Acid Injection, USP 12 mcg/2 mL 2 mL/vial

NDC NUMBER: 0517-6502-10

LOT NUMBER: 25090

EXPIRATION DATE: February 28, 2027