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 Lot 24394 of Cyanocobalamin Injection, USP manufactured and distributed by American Regent, Inc. is the subject of a recall. The recall of this product to CUSTOMER LEVEL was initiated due to glass particulate found in the reserved population of the batch.

Cyanocobalamin Injection, USP is a sterile solution intended for intramuscular or subcutaneous administration, indicated for the treatment of vitamin B12 deficiencies caused by malabsorption, pernicious anemia, and certain gastrointestinal disorders or surgeries.

Administration of Cyanocobalamin containing particulate matter may cause localized reactions, including irritation, pain, swelling, or granuloma formation. Given the intramuscular or subcutaneous routes of administration, when Cyanocobalamin Injection, USP is handled in accordance with the approved prescribing procedures, effects would be expected to remain confined to the injection site and are not anticipated to result in systemic adverse effects. The presence of the identified glass particle is considered to present a minimal, low toxicological risk following administration. As of October 7, 2026, American Regent has not received any reports of adverse events related to recalled batch, Lot 24394.

This recall is specific to the listed batch, Lot No. 24394. No other lots are impacted by this action.

PRODUCT: Cyanocobalamin Injection, USP 1000 mcg/mL, 1 mL/vial

NDC NUMBERS: 0517-0031-25 & 0517-0031-01

LOT NUMBER: 24394

EXPIRATION DATE: October 31, 2026