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 communicate that Lot(s) 25431 and 25431A of the product Cyanocobalamin Injection USP, manufactured and distributed by American Regent, Inc. is the subject of a recall to the RETAIL LEVEL. This recall associated with the batches listed was initiated due to a cellulose fiber found in the reserved population.

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 fiber particle is considered to present a minimal, low toxicological risk following administration. As of October 27, 2026, a review of the American Regent, Inc. safety database identified no serious adverse events associated with the recalled lot that could be attributed to the subject defect.

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

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

LOT NUMBERS: 25431 & 25431A

EXPIRATION DATE: 10/31/2027

FIRST DATE OF DISTRIBUTION: 5/20/2026