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 notice is to inform you of a product recall involving **Papaverine Hydrochloride, USP.**

Product: Papaverine Hydrochloride, USP

NDC No.: 0517-4002-25

Strength: 30 mg/mL

Package Size: 2 mL/vial

Manufactured By: American Regent, Inc.

Distributed By: American Regent, Inc.

Lot No: 25202; No other lots are impacted by this action.

Expiration Date: November 30, 2026

Recall of this product to the CONSUMER LEVEL if the product was administered to a patient.

The recall was initiated due to glass particles found in a reserve sample.

The administration of an injectable product containing particulate matter may result in local irritation or swelling. The particulate matter could travel through and block blood vessels in the heart, lungs or brain which can cause stroke and even lead to death. Patients with underlying cardiovascular or pulmonary compromise may be at increased risk. To date, American Regent, Inc. has not received any reports of adverse events related to this recall.