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.

Lot No. 24159N0C0 and 25193N0C0 of Levocarnitine 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 unlabeled vials, resulting in misbranded drug product due to the absence of required labeling information (e.g. product identification, lot number, expiration date).

Levocarnitine Injection, USP is indicated for the acute and chronic treatment of patients with an inborn error of metabolism which results in secondary carnitine deficiency and for the prevention and treatment of carnitine deficiency in patients with end stage renal disease who are undergoing dialysis.

PRODUCT INFORMATION	**Product**: Levocarnitine Injection, USP **Strength:** 200 mg/mL **Package Size:** 5 mL/vial **Manufactured By:** American Regent, Inc. **NDC No.**: 0517-1045-05, 0517-1045-01	**Lot No:** 24159N0C0 **Expiration Date:** June 2026 **Lot No:** 25193N0C0 **Expiration Date:** July 2027