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.

Sun Pharmaceutical Industries, Inc. has initiated a recall in response to an Out of Specification/Out of Tolerance results reported in unknown impurity in Related Substance test in Liothyronine Sodium Tablets 5 mcg/25 mcg, Batches DND0058A and DNC2204A at 18M long term condition, respectively. Sun Pharmaceutical Industries, Inc. initiated shipment of this product for Lot DNC2204A on February 21, 2022 and for Lot DND0058A on March 4, 2022.

| Product Name                               | Package Description Lot Number |          | NDC Number   | <b>Expiration Date</b> |
|--|--------------------------------|----------|--------------|------------------------|
| Liothyronine Sodium<br>Tablets, USP 5 mcg  | 100 tablets in<br>1 bottle     | DND0058A | 62756-589-88 | 12/2023                |
| Liothyronine Sodium<br>Tablets, USP 25 mcg | 100 tablets in<br>1 bottle     | DNC2204A | 62756-590-88 | 11/2023                |