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 this recall in response to an Out of Specification (OOS) result reported for individual unknown impurity in Liothyronine Sodium Tablets, 5 mcg. A total of thirteen (13) batches of Liothyronine Sodium Tablets, USP 5 mcg were Out of Specification.

Sun Pharmaceutical Industries, Inc. initiated shipment of this product on March 3, 2022.

Product Name	**Package Description**	**Lot Number**	**NDC Number**	**Expiration Date**
Liothyronine Sodium Tablets, USP 5 mcg	100 tablets in 1 bottle	DND0059A	62756-589-88	12/2023
Liothyronine Sodium Tablets, USP 5 mcg	100 tablets in 1 bottle	DND0060A	62756-589-88	12/2023
Liothyronine Sodium Tablets, USP 5 mcg	100 tablets in I bottle	DND0061A	62756-589-88	12/2023
Liothyronine Sodium Tablets, USP 5 mcg	100 tablets in 1 bottle	DND0062A	62756-589-88	01/2024
Liothyronine Sodium Tablets, USP 5 mcg	100 tablets in 1 bottle	DND0063A	62756-589-88	01/2024
Liothyronine Sodium Tablets, USP 5 mcg	100 tablets in 1 bottle	DND0064A	62756-589-88	01/2024
Liothyronine Sodium Tablets, USP 5 mcg	100 tablets in 1 bottle	DND0065A	62756-589-88	01/2024

Liothyronine Sodium Tablets, USP 5 mcg	100 tablets in 1 bottle	DND0180A	62756-589-88	01/2024
Liothyronine Sodium Tablets, USP 5 mcg	100 tablets in 1 bottle	DND0181A	62756-589-88	01/2024
Liothyronine Sodium Tablets, USP 5 mcg	100 tablets in 1 bottle	DND0182A	62756-589-88	01/2024
Liothyronine Sodium Tablets, USP 5 mcg	100 tablets in 1 bottle	DND0183A	62756-589-88	01/2024
Liothyronine Sodium Tablets, USP 5 mcg	100 tablets in 1 bottle	DND0184A	62756-589-88	01/2024
Liothyronine Sodium Tablets, USP 5 mcg	100 tablets in 1 bottle	DND0597A	62756-589-88	02/2024