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 inform you of a product recall by **Pfizer for Levoxyl (levothyroxine sodium tablets)** at the retail level.

Pfizer is initiating this recall due to confirmed out-of-specification (OOS) assay during stability testing. The overall potential risk to the patient arising from this issue is considered to be low. To date, Pfizer has not received any reports of adverse events associated with this issue.

Levothyroxine sodium tablets, USP)

NDC Number	Lot Number	Expiration Date	Strength	Configuration/Count
60793-851-01	24C11	FEB-2026	50 mcg	100 tablets per bottle