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.

A recall request of product has been initiated, Epinephrine/Lidocaine (0.25mg/mL, 7.5 mg/mL) Sterile Ophthalmic Injection (NDC NUMBER: 71384-0640-01) at our 503B facility, Imprimis NJOF, situated at 1705 US-46, Suite 6B, Ledgewood, NJ 07852. This recall is for two unexpired lots, 23JUL028 and 23AUG053, due to subpotent Epinephrine levels (<90%). This recall action has been prompted by a thorough investigation into a stability deviation, which conclusively established that the stability data for our Epi/Lido product does not support the assigned 360-Day expiration date. As a precaution, the assigned expiration date has been reduced to 180 days.

Lot Number	**Date Compounded On**	**Expiry Date**	**Quantity Released (20 units per box)**	**Quantity on Hand at Imprimis**
23JUL028	8/8/2023	8/1/2024	8120 units (406 bags)	0 units remaining
23AUG053	9/6/2023	8/30/2024	7740 units (387 bags)	0 units remaining