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.

Spectra Medical Devices, LLC has received the recall notification from the manufacturer, Huons about the product LIDOCAINE HYDROCHLORIDE INJECTION. Spectra is a distributor of this product.

Product Affected:

Strength	NDC Number	Dosage Unit	Configuration / Count
10 mg/mL	73293-0001-2	5 mLvial	10 vials in a carton
20 mg/mL	73293-0003-2	5 mL vial	10 vials in a carton
10 mg/mL	73293-0001-1	5 mL vial	5 ml single use vial
20 mg/mL	73293-0003-1	5 mL vial	5 ml single use vial