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.

Huons Co. Ltd. is recalling all lots of the below referenced product, **BUPIVACAINE HYDROCHLORIDE IN DEXTROSE INJECTION, USP, ALL LOTS.** Huons is initiating this recall due to quality issues identified during a recent FDA manufacturing facility inspection. We began shipping this product on or about **April 1, 2023.** Use of or consumption of this product may include a potential health hazard.

This recall should be carried out to the distributor, repackager, and user levels.

During this period, Huons has only received a small number of reports of drug ineffectiveness.

Strength	**NDC Number**	**Dosage** Unit	**Configuration/Count**
7.5 mg/ml	73293-0002-2	2ml ampule	50 ampules in 1 package
7.5 mg/ml	73293-0002-1	2ml ampule	Single-use Ampule label