

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.

Shenzhen Techdow Pharmaceutical Co., Ltd. is conducting a voluntary recall of the affected batch identified below due to a reassessment of the manufacturing deviation that may be deemed to have potential impact on sterility assurance.

This recall is being initiated as a precautionary measure to protect patient safety and with the knowledge of the US Food and Drug Administration.

No other batches from this product are being considered for market action at this moment. The information regarding the Batch is as follows:

Product: Heparin Sodium Injection, USP 5,000 USPU/mL, 1mL-multi dose vial

Batch/Lot Number: CU03232A

Quantity: 251,000 vials (in 10,040 packs)

NDC and package configuration: NDC 81952-111-06, 25 vials per carton

Expiration date: January 2028