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.

STAQ Pharma, Inc., Non-Resident Outsourcing Facility License No. NSF 143, is notifying you that STAQ Pharma, Inc. is initiating a recall on the hospital and surgery center level involving Ropivacaine Hydrochloride, 2 mg/mL, 500 mL in 500 mL IV Bags (NDC: 73177-0109-26). This recall is being implemented due to the sterile intravenous bag manufacturer incorrectly releasing and selling a lot of sterile, 500 mL IV bags to STAQ Pharma, Inc. that failed internal in-process quality control leak testing. Using a leaking IV bag could compromise sterility or can cause suboptimal infiltration therapy. The recall is being enacted with the knowledge of the Food and Drug Administration. One customer within the state of California received the recalled product.

The following product lots are affected:

Drug	NDC	Lot Number	Expiration Date
Ropivacaine Hydrochloride, 2 mg/mL, 500 mL in 500 mL IV Bag	73177-0109-26	23109472A	10 March 2024
		23109473A	11 March 2024
		23109474A	13 March 2024
		23109491A	16 March 2024
		23109492A	19 March 2024
		23109501A	25 March 2024
		23109520A	10 April 2024
		23109521A	13 April 2024
		23109522A	03 April 2024