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.

Hospira, Inc., a Pfizer company, is recalling the below-referenced lots of **ABBOJECT® products**; **4.2% Sodium Bicarbonate Injection, USP, 1% and 2% Lidocaine HCl Injection, USP** due to the possibility of glass particulates in the products. Pfizer completed a Health Hazard Assessment which concluded that the use of the impacted product has an unlikely probability of occurrence of adverse effects such as myocardial infarction, stroke, pulmonary embolism, deep vein thrombosis, and thrombophlebitis or mild pain. The potential risk to the patient arising from this issue is considered to be low.

To date, Pfizer has not received reports of any adverse events associated with this issue for these lots.

4.2% Sodium Bicarbonate Injection, USP

Product	NDC	Lot Number	Expiration Date	Strength	Configuration/Count
4.2% Sodium Bicarbonate Injection, USP Glass ABBOJECT® Syringe	Carton 0409-5534-24 Case 0409-5534-14	GJ5007	1AUG2024	5 mEq/10mL (0.5 mEq/mL)	1 vial and injector per carton; 10 cartons per bundle; Case Pack 5 X 10 – 10mL

1% Lidocaine HCl Injection, USP

Product	NDC	Lot Number	Expiration Date	Strength	Configuration/Count
1% Lidocaine HCl	Carton	42290DK	1JUN2024	50 mg /5mL	1 vial and injector per

Injection, USP	0409-4904-11			(10 mg/mL)	carton;
LIFESHIELD®	Case				10 cartons per bundle;
Glass ABBOJECT® Syringe	0409-4904-34				Case Pack 5 X 10 – 5mL

2% Lidocaine HCl Injection, USP

Product	NDC	Lot Number	Expiration Date	Strength	Configuration/Count
2% Lidocaine HCl	Carton	GH6567	1JUL2024	100 mg/5mL	1 vial and injector per
Injection, USP	0409-4903-11			(20 mg/mL)	carton;
LIFESHIELD®	Case				10 cartons per bundle;
Glass ABBOJECT® Syringe	0409-4903-34				Case Pack 5 X 10 – 5mL