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.

Fentanyl Citrate Injection, USP - CII

Tray NDC	Tray Lot Number	Vial NDC	Vial Lot Number	Expiration Date	Strength	Configuration/Count
0409-9094-22	13405DK	0409-9094-12	13-405-DK	1JUL2021	50 mcg/mL	Tray containing 25 x 2 mL Single-dose Fliptop Vials
0409-9094-22	17096DK	0409-9094-12	17-096-DK	1NOV2021	50 mcg/mL	Tray containing 25 x 2 mL Single-dose Fliptop Vials

Hospira Inc., a Pfizer company, is recalling the above referenced lots of **Fentanyl Citrate Injection, USP**. Pfizer initiated this recall due to confirmed customer reports for vials with the potential for loose metal overseal crimp defects. The recall of the above referenced lots of **Fentanyl Citrate Injection, USP** is being conducted to the **Hospital/Institution level**.