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.

PROPOFOL Injectable Emulsion, USP (contains Benzyl Alcohol)

Tray NDC	Vial NDC	Lot Number	Expiration Date	Strength	Configuration/Count
0409-4699-24	0409-4699-54	EA7470	01 JUN 2023	1 g/100 mL (10 mg/mL)	Tray of 10 x 100 mL Single Patient Use Fliptop Vials

Hospira Inc., a Pfizer company, is recalling the above referenced lot of **PROPOFOL Injectable Emulsion**. Pfizer initiated this recall due to a visible particulate observed in two vials during annual examination of retention samples. Pfizer completed a Health Hazard Assessment, which concluded that the use of the impacted product has a remote probability of being associated with potential adverse events, such as blockage of blood vessels, including decreased blood flow to the brain, heart attack, pulmonary embolus, and tissue necrosis. Hypersensitivity reactions and transmission of infectious disease can also occur. The overall potential risk to patients arising from this issue is considered to be medium.

The recall of the above referenced lot of **PROPOFOL Injectable Emulsion** is being conducted to the **User level**. The affected product lot was distributed in **July 2020**.