The Board of Pharmacy has received notice of the following product recall:

10% LMD in 5% Dextrose Injection (Dextran 40 in Dextrose Injection, USP)

NDC	**Lot Number**	**Expiration Date**	**Strength**	**Configuration/Count**
0409-7418-13 (Unit of Use) 0409-7418-03 (Unit of Sale)	87-095-JT	1MAR2020	10 g /100 mL	500 mL single-dose flexible containers; 12 overwrapped individual containers per case

Hospira, Inc., a Pfizer company, is recalling the above lot of **10% LMD in 5% Dextrose Injection** (Low Molecular Weight Dextran for Intravenous Administration **[Dextran 40 in Dextrose Injection, USP]**), to the hospital level, due to a manufacturing molding process defect resulting in variations on the additive port surface, which may lead to potential product leakage. The product was shipped product between **September 2018** and **July 2019**.

The Board of Pharmacy strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.