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.

Vancomycin Hydrochloride for Injection, USP

For intravenous use

1.5 g Single-dose Fliptop Vial, Sterile Powder

Carton NDC	Vial NDC	Lot Number	Expiration Date	Strength	Configuration/Count
0409-3515-01	0409-3515-11	33045BA	1SEP2023	1.5 g/vial	10 units/carton, 10 cartons/case

Hospira Inc., a Pfizer company, is recalling the above referenced lot of **Vancomycin Hydrochloride for Injection, USP,** to the **User level,** due to a confirmed report for the presence of glass fragments within a single vial. Pfizer completed a Health Hazard Assessment, which concluded the use of the impacted product has an unlikely probability of occurrence of limited severity adverse events such as phlebitis, end-organ granuloma or micro-embolic effects, or gastrointestinal trauma. The potential risk to a 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.