

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.

Zydus Pharmaceuticals (USA) Inc. is recalling the drug products mentioned below at the **HOSPITAL LEVEL:**

| <b>Product</b>                             | <b>NDC Number</b> | <b>Lot Number</b> | <b>Expiry Date</b> | <b>Pack Size</b>                    | <b>Distribution Start Date</b> | <b>Distribution End Date</b> |
|--|-------------------|-------------------|--------------------|-------------------------------------|--------------------------------|------------------------------|
| Acyclovir Sodium injection 50 mg/mL, 20 mL | 68382-0049-10     | L000155           | Dec 21             | 10x20 mL,<br>Single-Use Vial pack   | Start Dt.<br>12/23/2020        | End Dt.<br>01/05/2021        |
| Acyclovir Sodium injection 50 mg/mL, 20 mL | 68382-0049-10     | L000156           | Jan 22             | 10x20 mL,<br>Single –Dose Vial pack | Start Dt.<br>01/07/2021        | End Dt.<br>01/07/2021        |
| Acyclovir Sodium injection 50 mg/mL, 10 mL | 68382-0048-10     | L000126           | Dec 21             | 10x10 mL,<br>Single-Use Vial pack   | Start Dt.<br>12/23/2020        | End Dt.<br>01/05/2021        |
| Acyclovir Sodium injection 50 mg/mL, 10 mL | 68382-0048-10     | L000127           | Dec 21             | 10x10 mL,<br>Single-Use Vial pack   | Start Dt.<br>01/07/2021        | End Dt.<br>01/07/2021        |

Zydus Pharmaceuticals (USA) Inc. has initiated a recall of the drug products mentioned above after receiving three complaints of similar nature (“Crystallization in Vials”) in Acyclovir Sodium injection 50 mg/mL - both pack sizes of 10 mL and 20 mL.