

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 one lot of the drug product mentioned below at the **HOSPITAL LEVEL**:

Product	NDC Number	Lot Number	Expiry Date	Count	Distribution Start Date	Distribution End Date
Fulvestrant Injection 250mg/5mL, (50mg/mL)	70710- 1688-8	B200076	January 2024	250mg/5mL (50mg/mL), 5mL Pre- Filled Syringe	05/04/2022	06/13/2022

Zydus Pharmaceuticals (USA) Inc. has decided to initiate a recall of one lot of Fulvestrant Injection 250mg/5ml (50mg/mL) based on an out-of-specification (OOS) result observed during a stability sample test for Batch Number B200076 at the three months real-time data point. Our investigation of this OOS is currently under process but out of an abundance of caution and our continuous focus on patient safety, we are proactively recalling this referenced batch at the **Hospital Level**.