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.

This communication is to notify you that Zydus Pharmaceuticals (USA) Inc. (“Zydus”), is recalling the drug product mentioned below at the Retail Level.

This recall has been initiated due to a product complaint related to a single tablet having the foreign material embedded resembling a metal shaving. While the complaint is still under investigation, the recall is being carried out as a precautionary measure to ensure the patient safety. Consumption of an affected product may result in varying degrees of gastrointestinal irritation. The product was shipped to our customers in the month of May 2024.

PRODUCT: Venlafaxine Tablets, USP, 75mg 100’s Count Bottle Pack

NDC NUMBER: 68382-021-01

LOT NUMBER: M314265

EXPIRY DATE: 10/2025

DISTRIBUTION DATES: 5/1/2024 - 5/29/2024