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 is to inform you of a product recall being conducted by Bionpharma Inc. for one lot of Divalproex Sodium Delayed-Release Tablets USP, 500 mg distributed between February 17, 2026 and April 27, 2026. Bionpharma has initiated a recall of the impacted lot, and it is being conducted in cooperation with the Food and Drug Administration (FDA).

A single foreign tablet was found in one 500's count bottle of the below-referenced lot of the product. Based on the investigation, this is an isolated incident with an extremely low frequency of occurrence. The foreign tablet is readily distinguishable from Divalproex Sodium Delayed-Release Tablets USP, 500 mg based on their physical characteristics, including shape (round vs. oval), color (off-white vs. blue), size (10 mm vs 19 mm) and imprint (C2 vs 513). So, one product cannot be mistaken for another. However, out of an abundance of caution, Bionpharma has decided to recall the batch to the retail level.

PRODUCT: Divalproex Sodium Delayed-Release Tablets USP, 500 mg, 500's Count

NDC NUMBER: 69452-435-30

LOT NUMBER: 704C003

EXPIRATION DATE: 12/2028