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 Ergocalciferol Capsules USP, 1250 mcg (Vitamin D) distributed between 5-18-2026 and 8-28-2026. Bionpharma has initiated a recall of the impacted lot, and it is being conducted with the knowledge of the Food and Drug Administration (FDA).

A single foreign capsule was found in one 100'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 capsule is readily distinguishable from Ergocalciferol Capsules USP, 1250 mcg (Vitamin D) based on their physical characteristics, including color (white vs. green), size (19 mm vs 13 mm) and imprint (PA2120 vs PA140). 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.

It is important to note that Bionpharma has not received any adverse event reports to date relating to this issue and no other lot of Bionpharma's Ergocalciferol Capsules USP, 1250 mcg (Vitamin D) is impacted by this recall.

PRODUCT: Ergocalciferol Capsules USP, 1250 mcg (Vitamin D), 100's Count

NDC NUMBER: 69452-151-20

LOT NUMBER: S2642136

EXPIRATION DATE: DECEMBER 2028