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.

Sun Pharmaceutical Industries, Inc. would like to inform you of a product recall involving, Perampanel 6mg tablet, CIII.

This recall has been initiated in response to a report from pharmacist that Perampanel 10mg tablets were inside a bottle of Perampanel 6mg tablets and dispensed as Perampanel 6mg. The Pharmacy repackages Perampanel 6mg tablets into a blister pack and noted the color / strength difference after the repackaging of the subsequent shipment.

While Sun Pharma is not able to confirm that the Perampanel 10 mg tablets were labeled as Perampanel 6mg tablets as the bottle was discarded and the product was not directly distributed by Sun Pharma, the subject lot is being recalled in the event of a very low probability that Sun Pharma inadvertently labeled and distributed Perampanel 10 mg bottle(s) as Perampanel 6 mg. The tablets are embossed with the correct strength, therefore, visual identification of the strength is possible reducing the likelihood of consuming an incorrect dose.

Based on the Health Hazard Evaluation, ƒAlthough.the.recommended.daily.dose.can.be.up.to.78.mg?.unintended.higher.exposure.without.appropriate.titration.may.increase.the.risk.of.adverse.reactionsƒ

Sun Pharmaceutical Industries, Inc. initiated shipment of this product between November 26, 2026 and April 13, 2026.

PRODUCT: Perampanel 6mg tablet, 30 count

NDC NUMBER: 51672-4206-6

LOT NUMBER: AE01763

EXPIRATION DATE: 9/30/2027