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 Pharma is recalling the following product:

Product Name	Package Description	Lot Number	NDC Number	Expiration Date
Divalproex Sodium delayed-release tablets, USP 500mg	100 Count	HAC1312A	62756-798-88	05/2024

This recall has been initiated in response to an out of specification result observed during routine stability testing of dissolution test. A Health Hazard Evaluation conducted by Sun Pharma found that the out of specification result may result in lack of efficacy of batch HAC1312A.

Sun Pharmaceutical Industries Inc. initiated shipment of this product on 12/13/2021.

This recall should be carried out to the retail level.