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.

Product Name	Package Description	Lot Number	NDC Number	Expiration Date
Pregabalin Capsules, 50 mg	100 Count	DNC0432A	47335-687-88	01/2023

Sun Pharma is revising an earlier letter dated October 2 l, 2021 in regards to the same lot number of Pregabalin Capsule 50 mg in the above table, Lot DNC0432A. Per FDA classification dated December 22, 2021, **this recall is revised from the depth of wholesale level to retail leve**l.

This recall has been initiated in response to the Particle Size Distribution (PSD) and Bulk Density (BD) of the API used in Prcgabalin Capsules; 50mg was not meeting the specification limit. Sun Pharmaceutical Industries lnc. initiated shipment of this product on 07/08/2021.