The Board of Pharmacy has received notice of the following product recall:

NDC	Product Description	Lot Number	Exp Date
13107-0058-01	Acetaminophen and Codeine Tablets USP 300/15mg Tablets	058180015A	07/2021
		058180016A	07/2021
		05818017B1	09/2021
13107-0059-99	Acetaminophen and Codeine Tablets USP 300/30mg Tablets	059180023A	05/2021
		059180024A	05/2021
		059180025A	05/2021
13107-0060-01	Acetaminophen and Codeine Tablets USP 300/60mg Tablets	060180017A	06/2021
		060180019B1	06/2021
		060180020A	06/2021
		060180021A	06/2021

Aurobindo is recalling the product above because the subject batches were manufactured utilizing a new manufacturing process that was not adequately validated to the expectations of current best industry practices. Aurobindo began shipping these batches to customers nationwide July 11, 2018 through June 19, 2019.

The Board of Pharmacy strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.