

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

NDC	Product Description	Lot Number	Exp Date
13107-031-34	Mirtazapine Tablets USP 15mg	031180028A	03/2021

Aurobindo is recalling the product above because this batch was manufactured in a processing area in which water leakage was observed. Although there was no direct contact of any component with the subject water during the leakage events, the subject batches are being recalled out of an abundance of caution. No other Mirtazapine batches are affected. Aurobindo began shipping this batch to customers nationwide August 9, 2018 through April 11, 2019.

This recall is to the retail level.

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.