

*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.*

Zydus Pharmaceuticals (USA) Inc., is recalling two lots of the drug product mentioned below at the RETAIL LEVEL. Zydus Pharmaceuticals (USA) Inc. has decided to initiate a recall of two lots of Oxybutynin Chloride Extended-Release Tablets, USP 10 mg based on an out-of-specification (OOS) result observed during testing of one of the stability samples for lot number M300652 at the six months long term time point. Our investigation of this OOS is currently ongoing. However, out of an abundance of caution, and our continuous focus on patient safety, we are proactively recalling this referenced batch at the Retail Level.

<b>Product</b>	<b>NDC Number</b>	<b>Lot Number</b>	<b>Expiry Date</b>	<b>Count</b>	<b>Distribution Start Date</b>	<b>Distribution End Date</b>
Oxybutynin Chloride Extended-Release Tablets, USP 10 mg	68382-256-05	M300652	December 2024	500's count	06/28/2023	06/28/2023
Oxybutynin Chloride Extended-Release Tablets, USP 10 mg	68382-256-01	M300651	December 2024	100's count	06/14/2023	09/18/2023