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.

This is to inform you of a product recall involving Amlodipine and Olmesartan Medoxomil Tablets, 10 mg and 20 mg by Ascend Laboratories.

An out-of-specification (OOS) result was observed during the 12th month of dissolution test analysis for Olmesartan content (LT conditions: 25°C / 60% RH) in Amlodipine and Olmesartan Medoxomil Tablets, 10 mg/20 mg, batch number 25121311. Of important note, dissolution test analysis for amlodipine contents were found within specification. Amlodipine and Olmesartan Medoxomil is a combination drug. Amlodipine besylate is a calcium channel blocker and Olmesartan Medoxomil is an angiotensin II receptor blocker; in combination, this prescription drug is indicated for treatment of hypertension to lower blood pressure. Lowering blood pressure reduces the risk of fatal and nonfatal cardiovascular events. Lowering blood pressure reduces the risk of fatal and nonfatal cardiovascular agents, primarily strokes and myocardial infarctions. Delayed dissolution of the product may delay the onset of the drug's pharmacological action.

Our firm began shipping this product on August 11, 2025.

PRODUCT: Amlodipine and Olmesartan Medoxomil Tablets, 10 mg and 20 mg, 30 tablet bottles

NDC NUMBER: 67877-500-30

LOT NUMBER: 25121311

EXPIRATION DATE: March 31, 2028