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.

Ascend Laboratories, LLC would like to inform you of a product recall involving Amlodipine and Olmesartan Medoxomil Tablets, 5mg/40mg to the retail level. Please reference lot-specific information included below.

An out-of-specification (OOS) result was observed during dissolution testing for Olmesartan content of reserve samples for batch number 24123460 during stability testing at 12 months, 25°C/60%RH.

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. Delayed dissolution of the product may delay the onset of the drug's pharmacological action.

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

PRODUCT: Amlodipine and Olmesartan Medoxomil Tablets 5 mg/ 40 mg, 30 tablets/ bottle

NDC NUMBER: 67877-501-30

LOT NUMBER: 24123460

EXPIRATION DATE: October 31, 2027