

*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/40 mg by Ascend Laboratories.

During testing of the reserve sample for Amlodipine and Olmesartan Medoxomil Tablets 10 mg/40 mg (Batch No. 25120696), an out-of-specification (OOS) result was identified for the dissolution assay of the Olmesartan Medoxomil component. The dissolution results for Amlodipine were within the established acceptance criteria and met the applicable specification requirements.

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 June 4, 2025.

PRODUCT: Amlodipine and Olmesartan Medoxomil Tablets 10 mg/40 mg, 30 tablets per bottle

NDC NUMBER: 67877-502-30

LOT NUMBER: 25120696

EXPIRATION DATE: December 31, 2026