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.

Ajanta Pharma USA, Inc., would like to inform you of a product recall involving Fenofibrate Capsules USP 200 mg.

The recalled lot was distributed between 4/28/26 to 5/8/26 to wholesalers and distributors and drug chain stores nationwide. This recall should be carried out to the retail level.

This recall has been initiated as a precautionary measure due to concerns identified during the review of the dissolution investigation for the batch, PA02216. Although the batch met release specifications, we are recalling the product to ensure continued compliance with quality standards.

PRODUCT: Fenofibrate Capsules USP 200 mg, 100 count

NDC NUMBER: 27241-120-04

BATCH NUMBER: PA02216

EXPIRATION DATE: 12/2029

MANUFACTURER DATE: 01/2026