

*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 recall has been initiated due to a market complaint received concerning a stray Rasagiline Mesylate tablet discovered in an unopened stock bottle of Ibuprofen and Famotidine. Following internal investigation, the stray product was later identified as the 1 mg Rasagiline Mesylate tablets manufactured by Alkem Laboratories Limited. As a precautionary measure, the site has decided to recall the affected lot.

This recall is being conducted with the knowledge of the Food and Drug Administration. This recall should be conducted to the RETAIL level.

**PRODUCT NAME:** Ibuprofen and Famotidine Tablets, 800/26.6 mg

**LOT NUMBER:** 23140190

**NDC NUMBER:** 67877-626-90

**EXPIRATION DATE:** December 2024