

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.

Akron Pharma is recalling ONELAX DOCUSATE SODIUM LIQUID STOOL SOFTENER LAXATIVE (DOCUSATE SODIUM 50MG/5ML), NDC 71399-0039-06 specifically the following lot:

| <u>Lots</u> | <u>Expiration Date</u> | <u>SIZE</u> |
|-------------|------------------------|------------------|
| ODS230001A | 10/2025 | 16 FL OZ (473mL) |
| ODS230002A | 10/2025 | 16 FL OZ (473ml) |
| ODS230003A | 11/2025 | 16 FL OZ (473mL) |

This recall is at the retail level.

This recall has been initiated because the product is manufactured using an active pharmaceutical ingredient (API) produced by Badri Vishal Chemical Private Limited. Badri Vishal Chemical Private Limited is under an FDA import alert. We began shipping these lot on 04 MAR 24.

Immediately examine your inventory and quarantine product subject to recall.

Please note that all product subject to this recall should be quarantined, inventoried, and returned.

To arrange product returns, please notify Akron Pharma.