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.

LEO Pharma is writing to inform you of a U.S. recall of Adbry® (tralokinumab-ldrm) 300 mg/2mL Single-dose Autoinjector (Lot number 003E24A), due to the presence of particulate matter in one unit from the lot, which lab tests have identified as wool fiber. The lot was manufactured and packed for the U.S. market at the Catalent Indiana, LLC (Novo Nordisk) facility in Bloomington, Indiana, in September 2024.

The product lot was distributed nationwide to wholesalers and distributors in the United States from May 6, 2025 through June 10, 2025.

PRODUCT: Adbry® (tralokinumab-ldrm) 300 mg/2mL Single-dose Autoinjector

NDC NUMBER: 50222-350-02

LOT NUMBER: 003E24A

EXPIRATION DATE: 04/30/27