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 that Unichem Pharmaceuticals (USA), Inc. is recalling Amiodarone Hydrochloride Tablets USP 200 mg, 60 count bottles Lot ZMRH25054, due to a confirmed product complaint of a metal fragment embedded in a tablet. discovered by a patient after splitting the tablet**.**

This recall is being made with the full knowledge of the United States Food and Drug Administration and should be carried out as a Retail Level Recall. Unichem has not received any reports of any adverse events related to this recall to date. Should you need to report an adverse event please call: +1 (866) 562-4616.

Amiodarone Hydrochloride Tablets USP, 200 mg, are white to off-white, round, flat-faced tablets with beveled edges. debossed with "U" above a bisect on one side and "359" on the other side. If swallowed. a metal fragment may cause medically significant gastrointestinal irritation or injury, including mucosal laceration, bleeding, or localized tissue damage.

Unichem Pharmaceuticals (USA), Inc. distributed 23,850 bottles of this lot to US customers through our downstream trading partners from March 20, 2026, through April 15, 2026.

PRODUCT: Amiodarone Hydrochloride Tablets, USP 200 mg, 60 count bottles

NDC NUMBER: 29300-359-16

LOT NUMBER: ZMRH25054

EXPIRATION DATE: 07/2028