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.

AvKARE was notified that Amneal Pharmaceuticals LLC is initiating a Recall to the RETAIL LEVEL for specific batches of Chlorpromazine Hydrochloride Tablets, USP (50 mg and 100 mg) associated with a specific lot of auxiliary polyester coil, due to the detection of a micro-organism. No micro-organism was detected on any tablets from any of the retain samples tested from the batches being recalled. No market complaint or adverse event has been received for any of the recall specific batches for any microbial related issue. Out of an abundance of caution and commitment to quality Amneal initiated a recall.

This recall is being carried out with the knowledge of the US Food and Drug Administration. Further use of this product should immediately cease. The below marketed product is subject to the recall. This recall is to the Retail Level Only.

Product Description	**NDC**	**Initial Ship Dates**	**Lot & Expiration**
Chlorpromazine Hydrochloride Tablets, USP 50 mg	**50268-164-15**	**8/5/2024**	**Lot 46824 Exp. 09/30/2025**
		12/16/24	**Lot 47171 Exp. 12/31/2025**
Chlorpromazine Hydrochloride Tablets, USP 100 mg	**50268-165-15**	**10/28/24**	**Lot 47089 Exp. 12/31/2025**
		3/13/25	**Lot 47604 Exp. 3/31/2026**