

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 communication is to notify you that Zydus Pharmaceuticals (USA) Inc. (“Zydus”), is recalling the drug product mentioned in the table below at the Retail Level.

Zydus Pharmaceuticals (USA) Inc. is recalling Hydrocortisone Sodium Succinate for Injection, USP 100 mg/vial, lots listed in Table 1 below.

The recall is based on an Out of Specification (OOS) result for “Any Unspecified Degradation Product” detected in the product at the 4-month interval, during long-term and accelerated stability condition samples testing. The presence of this impurity above acceptable limits may present a potential health concern; however, the risk from exposure is unknown.

The product was shipped to our customers from June 2026 to September 2026.

Product	Vial NDC Number	Carton NDC Number	Lot Number	Expiry Date	Count	Distribution Start Date	Distribution End Date
Hydrocortisone Sodium Succinate for Injection, USP 100 mg/vial	70710-1907-1	70710-1907-1	LB00047A	03/2028	Single-dose Vials	Jun-10-2026	Aug-13-2026
	70710-1907-1	70710-1907-1	LB00048A	03/2028	Single-dose Vial	Aug-13-2026	Sep-17-2026