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.

Amneal Pharmaceuticals LLC is initiating this recall of 3 lots of Phytonadione Injectable Emulsion, USP 1mg/0.5mL (NDC 70121-1682-7) due to the result for the phytonadione photodegradant in related compounds testing exceeded the specification.

Phytonadione injectable emulsion is indicated for the treatment of coagulation disorders due to vitamin K deficiency or interference with vitamin K activity and for prophylaxis and treatment of vitamin K deficiency bleeding in neonates.

A potential reduction in therapeutic efficacy may occur however, this is a known product degradant with a low toxicological risk at the detected level.

Only the 3 lot numbers listed in the table below are being recalled.

NDC NUMBER	LOT NUMBER	EXPIRATION DATE	1ST SHIP DATE	LAST SHIP DATE
70121-1682-7	AP250506	04/2027	03/04/2026	05/04/2026
70121-1682-7	AP250521	05/2027	04/30/2026	06/15/2026
70121-1682-7	AP250522	05/2027	04/03/2026	06/08/2026