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 notice is to inform you of a product recall involving EMRYDI® RTU (**Pemetrexed Injection) 100 mg/10 mL and 500 mg/50mL.**

This recall is being carried out to the Hospital Level.

This voluntary recall is for two lots of PEMRYDI® RTU (**Pemetrexed Injection)** 100 mg/10 mL, (NDC # 70121-2453-1) and two lots of PEMRYDI® RTU (**Pemetrexed Injection)** 500 mg/50 mL, (NDC # 70121-2461-1) due to product discoloration observed in some vials.

This recall is being conducted following the receipt of market complaints reporting green to dark green discoloration in certain vials, resulting in product appearance outside the approved specification of **Clear, Colorless to pale yellow to greenish yellow solution**.

To date, no adverse events have been reported that have been attributed to the observed discoloration.

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

NDC Code	Strength	Lot number	Expiration Date	1st Ship Date
70121-2453-1	100mg/ 10mL	P400172	09/2026	02/11/2025
70121-2453-1	100mg/ 10mL	PA00050A	09/2027	12/18/2025
70121-2461-1	500mg/ 50mL	P400157	08/2026	03/24/2025
70121-2461-1	500mg/ 50mL	PA00047A	09/2027	12/09/2025