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.

Mylan Institutional LLC (a Viatris company) is conducting a voluntary recall at the retail level of the below-listed lots of Mycophenolate Mofetil for Injection USP, 500 mg/vial, packaged in a carton containing 4 vials.

These lots are being recalled out of an abundance of caution following a report of a cloudy solution with undissolved matter present in the vial after reconstitution of the lyophilized cake.

These lots were distributed in the United States between 11/18/2025 to 06/24/2026.

To date, no reports of adverse events associated with this lot have been received. The risk associated with this event is considered to be Medium.

Mycophenolate mofetil is indicated for the prophylaxis of organ rejection in adult and pediatric recipients ≥ 3 months of age) of allogeneic kidney, heart, or liver transplants, in combination with other immunosuppressive agents.

PRODUCT: Mycophenolate Mofetil for Injection USP, 500 mg/vial, carton of four vials

NDC NUMBER: Carton- 67457-386-81

Vial- 67457-386-00

LOT NUMBER	EXPIRATION DATE
3242616	April 2027
8220216	Nov 2027