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 recall at the retail level of the below-listed lot of Mycophenolate Mofetil for Injection USP, 500 mg/vial, packaged in a carton containing 4 vials.

This lot is 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.

This lot was distributed in the United States between 09/23/2025 and 10/28/2025.

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

NDC NUMBER: 67457-386-81 [Carton Label] 67457-386-00 [Vial Label]

LOT NUMBER: 3241653

EXPIRATION DATE: April 2027