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 is to inform you that Sandoz Inc. ("Sandoz") has initiated a product withdrawal related to the incorrect Country of Origin printed on the drug product label.

The label on the drug product bottle for Mycophenolate Mofetil, USP 250 mg capsules (100 count), Mycophenolate Mofetil, USP 500 mg tablets (100 count) and Mycophenolate Mofetil, USP 500 mg tablets (500 count) states Product of Slovenia instead of the required Product of India.

Product Name	**NDC Number**	**Lot Number**	**Expiration Date**
Mycophenolate Mofetil, USP 250 mg Capsules	0781-2067-01	PE9742	10/2026
Mycophenolate Mofetil, USP 500 mg Tablets	0781-5175-01	PE9749 PG7363 PG7362 PH3159	10/2026 10/2026 11/2026 12/2026
Mycophenolate Mofetil, USP 500 mg Tablets	0781-5175-05	PJ0703	12/2026