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.

Micafungin for Injection, 50 mg and 100 mg lyophilized vials

Product	NDC No.	Batch No.	Expiration Date	Configuration
Micafungin for Injection, 50 mg vials	70594-036-01	467111	01/2023	Single vials in indivdual cartons; 32 vials/case
Micafungin for Injection, 10 mg vials	70594-03 7-01	467129	01/2023	

Xellia Pharmaceuticals USA LLC is recalling the above referenced batches of Micafungin for Injection, 50mg and 100mg lyophilized vials. This recall is being performed due to the omission of certain safety information from the package insert. This includes aspects of Adverse Reactions, Drug Interactions, and Use in Specific Populations. While there are no issues with the quality of the drug product, complete prescribing information is required for safe and effective use of this injectable prescription product.

This recall is being performed to the retail/pharmacy level. This product was shipped between June 8, 2021, and July 20, 2021