

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 of a product recall involving Lisdexamfetamine Dimesylate Capsules, 40mg, 50 mg, 60 mg, and 70mg by Sun Pharmaceuticals Industries, Inc.

This market withdrawal has been initiated in response to Out of Specification (OOS) results observed in dissolution test for Lisdexamfetamine Dimesylate Capsules, 60 mg, Batch AD42648, during analysis at 9 month long term stability station (25°C, 60%RH). Additionally, three (3) other batches with the same common blend are impacted and, therefore, subject to market withdrawal.

Sun Pharmaceutical Industries, Inc. initiated shipment of this product on May 1, 2024.

Product Name	Package Description	Lot Number	NDC Number	Expiration Date
Lisdexamfetamine Dimesylate Capsules, 40 mg	100 count bottle	AD42646	57664-049-88	2/28/2026
Lisdexamfetamine Dimesylate Capsules, 50 mg	100 count bottle	AD42647	57664-050-88	2/28/2026
Lisdexamfetamine Dimesylate Capsules, 60 mg	100 count bottle	AD42648	57664-051-88	02/28/2026

Lisdexamfetamine
Dimesylate Capsules, 70
mg

100 count
bottle

AD42649

57664-052-
88

2/28/2026