

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.

Hikma Pharmaceuticals USA Inc. and its affiliates ("**Hikma**") are voluntarily initiating a market withdrawal of 18 lots of **Abirtega™ (abiraterone acetate tablets, USP), 250 mg** and **Abiraterone Acetate Tablets, USP, 250 mg**.

This market withdrawal is being conducted as a precautionary measure. These lots were produced in the same manufacturing campaign as a lot that generated an out of specification result for weight variation. Testing parameters for these lots are within specification.

Abirtega™ (abiraterone acetate tablets, USP), 250 mg		Abiraterone Acetate Tablets, USP, 250mg	
NDC No. 82249-011-12		NDC No. 0143-9597-21	
120 count bottle		120 count bottle	
Lot Number	Expiry Date	Lot Number	Expiry Date
3046	July 2027	3053	August 2027
3048	August 2027	3054	August 2027
3049	August 2027	3055	August 2027
3050	August 2027	3056	August 2027
3051	August 2027	3057	August 2027
3052	August 2027	3058	August 2027
		3059	August 2027
		3060	August 2027
		3061	August 2027
		3062	August 2027
		3063	August 2027
		3064	August 2027