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.

Description	Lot #	Exp Date	NDC	UPC
ORPHENAD ER TAB 100MG SAN 100@	LA9243	11/30/2023	00185002201	30185002201
	LA7704	10/31/2023		
	LA7703	10/31/2023		
	KM0072	03/31/2023		
	KL3199	03/31/2023		
	KE7169	11/30/2022		
	KE4349	11/30/2022		
	KE4348	11/30/2022		
	KC3303	08/31/2022		
	KC0723	08/31/2022		
	JX6413	05/31/2022		
	JX6411	05/31/2022		
ORPHENAD ER TB 100MG SAN 1000	KS3939	03/31/2023	00185002210	30185002210

Sandoz is recalling the above item(s)/lot(s) due to an impurity that has the potential to be above FDA's acceptable daily intake. This recall is to the consumer level. Affected product started shipping August 2019.