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.

SpecGx LLC ("SpecGx"), a Par Health company, is recalling four (4) lots of Oxycodone and Acetaminophen Tablets USP 7.5 mg/325 mg and 10 mg/325 mg at the retail level.

SpecGx has initiated a recall because of the potential for the debossing to be missing or partially missing on one side of the tablet. Missing debossing is unlikely to cause an adverse event. Only the products and lots listed in the table above is impacted by the recall. This recall is being initiated with the knowledge of the Food and Drug Administration (FDA).

NDC	Product	Lot	Expiration Date
0406-0522-01	Oxycodone and Acetaminophen Tablets USP 7.5 mg/325 mg	0522J23493	03/2027
0406-0523-01	Oxycodone and Acetaminophen Tablets USP 10 mg/325 mg	0523J23904	05/2027
		0523J24426	06/2027
		0523J24427	06/2027