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
ACETA OS UD325MG/10.15ML P/D30	503715	12/31/2022	68094033062
	503693	12/31/2022	
	503679	12/31/2022	
ACETA OS325MG/10.15MLP/D UD100	503715	12/31/2022	68094033061
	503693	12/31/2022	
	503679	12/31/2022	
Acetaminophen Oral Suspension, 325 MG/10.15ML (Individual Cup)	503715	12/31/2022	68094033059
	503693	12/31/2022	
	503679	12/31/2022	
ACETAM CHL OS CUP 5ML PD UD100	503670	12/31/2022	68094023161
ACETAM CHL OS CUP 5ML PD UD30	503670	12/31/2022	68094023162
Acetaminophen Oral Suspension, 160 MG/5ML (Individual Cup)	503670	12/31/2022	68094023159

Precision Dose is recalling the above items/lots in response to the recall by the manufacturer (Perrigo) which included the above affected lots that were repackaged by Precision Dose. Perrigo notified Precision Dose that one of their suppliers had an out of specification result of Total Aerobic Microbial Count (TAMC) affecting a single drum of raw material. This recall is to the retail level. Affected product started shipping October 8, 2021.