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 recall has been initiated due to a notification received through Prodigy Health which is an Accord Healthcare distributor based on a recall (Distributor level) initiated by Accord Healthcare, US subsidiary of Intas Pharma the manufacturer of the phenylephrine finished dose vials used to produce these lots of phenylephrine vials. Reference recall titled "Voluntary recall of Drug Products manufactured at FEI # 3004011473, Intas Pharmaceuticals Ltd." that was initiated on February 7, 2023 by Accord Healthcare, Inc., due to an FDA inspection that was conducted at the Intas Pharma Gujarat, India facility with several Observations on Form 483.

Product Information:

F3001	71449-001-11	Phenylephrine HCl 100 mcg/mL, 5 mL in 5 mL Syringe
F3002	71449-001-15	Phenylephrine HCl 100 mcg/mL, 10 mL in 10 mL Syringe
	71449-150-82 e 250 mL IV Bag	Phenylephrine HCl 40 mg (160 mcg/mL) (from FDP) added to 0.9% Sodium
	71449-148-94 e 250 mL IV Bag	Phenylephrine HCl 20 mg (80 mcg/mL) (from FDP) added to 0.9% Sodium

F3002 2230895 3/5/2023 F3002 2230911 3/11/2023	
F3002 2230911 3/11/2023	
F3002 2230913 3/18/2023	
F3001 2230960 3/12/2023	
F3002 2230994 3/27/2023	
F3002 2231006 4/1/2023	
F3352 2231017 3/12/2023	
F3360 2231026 2/23/2023	
F3002 2231030 4/12/2023	
F3360 2231051 3/11/2023	
F3001 2231080 4/9/2023	
F3002 2231109 4/19/2023	
F3002 2231126 5/6/2023	
F3002 2231134 5/10/2023	
F3002 2231140 5/14/2023	
F3002 2231142 5/20/2023	

F3002	2231156	5/29/2023
F3360	2231163	3/26/2023
F3352	2231199	4/30/2023
F3002	2231273	6/3/2023
F3002	2231285	6/10/2023
F3002	2231299	6/17/2023
F3360	2231304	5/7/2023
F3360	2231308	5/11/2023
F3002	2231331	6/26/2023
F3002	2330014	7/9/2023
F3002	2330025	7/15/2023