The Board of Pharmacy has received notice of the following product recall:

LEVEL OF NOTIFICATION: retail pharmacy

SUPPLIER: Sandoz

Description	Lot #	Exp Date	NDC	UPC
GATIFLOX OPH .5% SAN 2.5ML	290632F 10/31/20; 289210F 10/31/19		61314067225	36131467225
NEOMY+POLY B+DEX O/S FAL 5ML@	295342F 08/31/20; 287880F 10/31/19; 287881F 10/31/19; 290555F 11/30/19; 290556F 03/31/20; 290557F 03/31/20; 293386F 03/31/20; 293387F 03/31/20; 293388F 05/31/20; 293389F 02/29/20; 293390F 05/31/20; 293392F 07/31/20; 304966F 04/30/21; 295344F 10/31/20; 295345F 10/31/20; 295346F 10/31/20; 295347F 11/30/20; 298823F 08/31/20; 298825F 12/31/20; 304495F 01/31/21; 304496F 11/30/20; 304497F 01/31/21; 304963F 03/31/21; 304964F 05/31/21; 293391F 06/30/20		61314063006	36131463006

Sandoz is voluntarily recalling the above items and lots because certain safety label changes were not updated in the patient insert. This recall is to the pharmacy level. Affected product started shipping December 7, 2017.

The Board of Pharmacy strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.