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

PRODUCT DESCRIPTION: BD Microtainer® Tube w/ BD MicrogardTM Closure. K2EDTA additive. Lavender closure 250-500uL fill volume.

CAT #: 365974

Lot Number	UDI (GTIN, DI+PI)	Exp. Date
9017591	(01)30382903659747 (17)200630(10)9017591(30)50 (01)50382903659741 (17)200630(10)9017591(30)200	2020-06-30
9017593	(01)30382903659747 (17)200630(10)9017593(30)50 (01)50382903659741 (17)200630(10)9017593(30)200	2020-06-30
9052823	(01)30382903659747 (17)200731(10)9052823(30)50 (01)50382903659741 (17)200731(10)9052823(30)200	2020-07-31

BD is conducting a medical device recall for BD Microtainer® Tubes w/ BD MicrogardTM Closure, K2EDTA additive, Catalog# 365974. The above-referenced lots have been confirmed to have reduced or no additive within the tube reservoir. BD distributed affected lots between February 4, 2019 and April 5, 2019.

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.