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

Product Name	**Cat. No. (Ref)**	**Recall Date**	**UDI**	**Lot No.**	**Exp. Date**
BD Vacutainer® Eclipse™ Blood Collection Needle 22Gx1.25”	368608	**December 4, 2019**	**(17)231231(10)9010765(30)480 (01)50382903686082**	**9010765**	**December 31, 2023**
		July 17, 2019	(17)240131(10)9025826(30)480 (01)50382903686082	9025826	January 31, 2024
			(17)231231(10)8354527(30)480 (01)50382903686082	8354527	December 31, 2023
		March 7, 2019	(17)230731(10)8207894(30)480 (01)50382903686082	8207894	July 31, 2023

BD issued recall notices for BD Vacutainer® Eclipse™ Blood Collection Needle 22Gx1.25” for lot 8207894 dated March 7, 2019, and for lots 9025826 and 8354527 dated July 17, 2019.

BD has confirmed that one additional lot, 9010765, was found to be missing the bevel from the non-patient (NP) needle end of the Eclipse™ Blood Collection Needle. This NP needle is housed in a sleeve that encloses the needle after puncture to prevent any leakage. If there is no bevel, the needle may damage the sleeve, causing leakage. Additionally, it may make it difficult to insert the BD Vacutainer® blood collection tubes onto the NP end.

Distribution of lot 9010765 occurred beginning on February 1, 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.