

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

Product Description	Catalog No.	Lot No.	UDI	Expiration Date
BD PrecisionGlide™ Needle 19GX1-1/2IN	305187	9030812	30382903051879	March 2024

BD is initiating a recall for the one lot of BD PrecisionGlide™ Needle referenced in the table above because a portion of the lot was packaged with the incorrect size needle (22GX1-1/4IN). BD has not received any reports of related adverse events, and the anticipated health risk is low as it is expected the clinician will identify that the needle size is incorrect prior to use. BD distributed the affected lot between April-October 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.