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.

Product Name	**UDI**	**Catalog (Ref) No.**	**Lot No.**	**Expiration Date**	**Product Package Size**
BD Nexiva™ 20 GA 1.00 (1.1 x 25mm) Dual Port	382903835362	383536	1193055 1188953 1166273	6/30/2024	80

BD is conducting a medical device recall of three lots of the BD Nexiva ™ Closed IV Catheter System referenced in the table above. BD has confirmed through internal leak testing a breach in the product packaging. This breach in the product packaging renders the enclosed IV catheters non-sterile. If the issue is not identified prior to using the device, this poses an increased risk of introduction of microbes into the blood stream. The health effects range from no clinical effect up to bloodstream infection which may be life threatening.