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.

BD was notified by the manufacturer Fresenius Kabi USA, LLC ("Fresenius Kabi") that they are recalling the below-mentioned batches of 0.9% Sodium Chloride Injection, USP products in Single Dose freeflex® bags.

Fresenius Kabi has decided to take this action out of an abundance of caution due to customers reporting blue Break-off-Parts (BOPs) that have detached from the infusion port during manipulation.

Fresenius Kabi has conducted a Health Hazard Evaluation and concluded there is low risk for patients and healthcare professionals. For these batch numbers, no adverse drug reaction or harm has been reported.

Catalog Number/SKU	**Product Name**	**Lot Number**
1727170102	0.9% Sodium Chloride Injection, USP 50mL	6402481 6402482 6402428
1727170103	0.9% Sodium Chloride Injection, USP 100mL	6402153 6402297 6402298 6402377 6402378 6402379 6402429 6402430 6402431 6402432 6402433 6402434 6402512 6402574 6402576
1727170105	0.9% Sodium Chloride Injection, USP 250mL	6402290 6402291 6402411 6402412 6402419 6402424 6402425 6402426 6402427 6402479 6402480

1727170107	0.9% Sodium Chloride Injection, USP 1000mL	6402517 6402518 6402571 24EU10010