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.

| **Catalog No.** | **Recalled Product** |
|---|---|
| 260400 | ChloraPrep® One-Step 3 mL Applicator - Clear |
| 260415 | ChloraPrep® With Tint 3 mL Applicator - Hi-Lite Orange |
| 930400 | BD ChloraPrepTM Clear 3 mL Applicator |
| 930415 | BD ChloraPrepTM Hi-Lite OrangeTM 3 mL Applicator |

On June 23rd, 2020, Becton Dickinson and Company (BD) recalled the ChloraPrep™ 3mL product for those regions of the world where the product could be exposed to 30°C / 75%RH continuously for more than six months. The recall did not affect the mainland US, but after further consultation with the FDA and out of abundance of caution, BD has expanded the recall of the ChloraPrep™ 3mL products with catalog numbers listed in the table above in all 50 U.S. states. There has been no identified risk associated with the sterile ChloraPrep™ antiseptic solution within the applicator. To date, no complaints, adverse events, injuries, or deaths have been reported related to this recall.

This recall should be carried out to the user level.

The recall **does not** impact ChloraPrep™ 3mL products that are purchased in the bulk non-sterile configuration for kits.