

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.

Molded Products, Inc. is conducting a medical device recall of the MPC-130 See Luer Cap Set based on a confirmed complaint of the threaded sleeve not being engaged and becoming un-attached. Some units of sale packages (bags of 100 sets) were found to have sleeves that were unattached to the luer pin. The case carton and individual polybags are correctly labeled as Molded Products' *MPC-130 Luer Lock Set, Lot# 20389*. The MPC-130 See Luer Cap Set contains (2) non-injectable end caps used to cap central venous catheters, Estimated % of lot affected: Less than 3.18%.

While no health risk has been identified, we have determined that certain units may not meet our quality standards due to a malfunction that affects product integrity. The condition may result in the device not remaining intact as the white protective cap is removed. Based on the nature of the defect, it is expected healthcare professionals would identify the nonconformance during routine inspection and discard the affected unit prior to patient use, preventing any patient contact. The primary impact is limited to inconvenience.

Distribution of the affected lot occurred from April 25, 2025, through August 27, 2025.

PRODUCT: See Luer Cap Set

UDI: +B144MPC1300/\$\$529105203895

LOT NUMBER: 20389

CATALOG NUMBER: MPC-130

EXPIRATION DATE: 04/15/2029