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.

ICU Medical, Inc. is issuing a recall to the user level, for a MISLABELLED lot of POTASSIUM CHLORIDE Inj. 20 mEq, NDC 0990-7075-26. The OVERWRAP label of lot 1023172, Expiration Date: 01-31-2026, incorrectly identifies the product as POTASSIUM CHLORIDE Inj. 10 mEq with NDC 0990-7074-26. The dosage is correctly printed on the labeling affixed to the product bag which is not visible when the 10 mEq OVERWRAP is in place.

If the incorrect dosage on 10 mEq overwrap is used instead of the correct 20mEq dosage printed on the product, an overdose of potassium chloride is possible. Overdose of potassium chloride can lead to hyperkalemia. ICU Medical has not received reports of adverse events associated with this issue to date.

The affected product lot was manufactured on 24 September 2024 and distributed in the United States between 15 November 2024 through 17 December 2024.

PRODUCT: POTASSIUM CHLORIDE Inj. 20 mEq - 100 mL Flexible Container

NDC NUMBER: 0990-7075-26

LOT NUMBER: 1023172

EXPIRATION DATE: 1/31/2026

LIST NUMBER: 070750453