

*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.*

This is to inform you of a product recall from Dr. Reddy's Laboratories Inc. involving Succinylcholine Chloride Injection, USP 200 mg/10 ml (20 mg-mL).

This recall is being initiated in reference to an observed Out-of-Specification (OOS) result during the 6-month stability testing. The test parameter, Methyl Paraben Content by HPLC, yielded an average result of 89.5%, which falls below the specification limit of not less than 90.0% and not more than 110.0%, and is therefore considered OOS. This recall is being carried out to the retail level.

A decrease in preservative concentration increases the risk of reduced sterility assurance. In this case, an out-of-specification (OOS) result was observed at 89.5%, falling below the specification range of 90.0% to 110.0%. Although the reduction in methylparaben content is minimal (0.5%), it could potentially impact the preservative system's effectiveness. Due to the absence of microbial testing data for the affected batch, a comprehensive assessment cannot be made. However, no adverse events or sterility-related issues have been reported to date for this batch.

The product Distribution Dates: February 18, 2025 - May 20, 2025.

PRODUCT: Succinylcholine Chloride Injection, USP 200 mg/10 ml (20 mg-ml) vial, 25 count

NDC NUMBER: 43598-666-25

LOT NUMBER: K250048

EXPIRATION DATE: 04/2026