B. Braun Medical – Recall of sodium chloride injection

- On August 2, 2023, B. Braun Medical announced a consumer-level recall of one lot of sodium chloride 0.9% injection due to a portion of the label to be missing or partially printed.
    - Other sodium chloride injection products that are not being recalled are available for patients to use
- The recalled lots were distributed between December 2022 and February 2023.

Product Description	NDC#	Lot # (Expiration Date)
0.9% sodium chloride injection in EXCEL plus IV container, 1000 mL	0264-5802-00	0061852531 (2/28/2025)

- Sodium chloride injection is indicated for:
    - Use in adults and pediatric patients as sources of electrolytes and water for hydration
    - Extracellular fluid replacement, treatment of metabolic alkalosis in the presence of fluid loss and mild sodium depletion
    - Use as a priming solution in hemodialysis procedures and may be used to initiate and terminate blood transfusions without hemolyzing red blood cells
    - Use as pharmaceutic aids and diluents for the infusion of compatible drug additives.
- Per B. Braun Medical, the leftmost portion of the container labeling was not printed and therefore information including the description, warnings, storage information and instructions for use were either missing or partially printed.
- Additionally, if the clinician notices the missing label, a delay in treatment may occur in order to obtain a new bag. If the clinician does not notice a portion of the container labeling is missing, and believes the container contains hypertonic solution, the solution administered for therapy will not have the desired effect and additional medical intervention may be required.
- Anyone with an existing inventory of the recalled product should stop use, distribution and quarantine the product immediately.
- For any questions regarding this recall, contact B. Braun Medical by phone at **1-833-425-1464**.