Baxter – Recall of 0.9% sodium chloride injection and 5% dextrose injection

- On July 6, 2017, Baxter announced a consumer-level recall of one lot of 5% dextrose injection and 4 lots of 0.9% sodium chloride injection due to the potential presence of leaks.

- A leak of the solution bag may allow for delay or interruption of therapy, under-delivery, unintended drug exposure, and microbial contamination. If not detected, the use of a solution bag with a leak could lead to a bloodstream infection or other serious adverse health consequences.

    — To date, there have been no reports of adverse events associated with this issue for leaks.

- The recalled lots were distributed between April 4, 2017 and June 1, 2017.

Product Description	NDC #	Lot # (Expiration date)
5% Dextrose Injection, 100mL VIAFLEX Plastic Container Multi Pack	00338-0017-38	P361618 (9/30/2018)
0.9% Sodium Chloride Injection, 100 mL VIAFLEX Plastic Container	00338-0049-38	P361501 (9/30/2018), P361667 (9/30/2018), P361790 (9/30/2018)
0.9% Sodium Chloride Injection, 250 mL VIAFLEX Plastic Container	00338-0049-02	Y229153 (9/30/2018)

- Dextrose injection is indicated as a source of water and calories.

- Sodium chloride injection is indicated as a source of water and electrolytes and for use as a priming solution in hemodialysis procedures.

- Anyone with an existing inventory of the recalled lots should stop use and distribution, and return all recalled product.

- For questions about this recall, please contact the Baxter Healthcare Center for Service at **1-888-229-0001**.