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.

The purpose of this is to inform you that B. Braun Medical Inc. (BBMI) is issuing a pharmaceutical recall for products L7500, LAC RNG INJ 1L, and L8000, NACL INJ 0.9% 1000ML due to the potential of fluid leakage at one of the weld sites.

The depth of this recall is to the hospital/healthcare facility level. This recall is being conducted with the knowledge of the United States Food & Drug Administration.

BBMI identified during a manufacturing in-process inspection that there is a potential for fluid leakage. It was determined that misalignment of the hole punch resulted in poor welds at the saddle weld location.

To date there have been no reports of serious injury or death associated with the issue. Product leakage can result in the potential delay of treatment while a new IV bag is obtained. If the bag contains hazardous medication, there is a potential for leakage of that medication with limited to significant exposure to hazardous drugs that may result in mild to severe injury to the patient, clinician, or other. If the leakage is not evident, the contents of the container may be contaminated due to compromising the sterile barrier of the product, potentially resulting in bloodstream infection. In some patients, bloodstream infection or delay in critical medication treatment may cause death.

Product Catalog Number	**NDC Number**	**Product Description**	**Expiration Date**	**Region Distributed**	**Lot Number**	**Distribution Range**
L7500	0264-7750-00	LAC RNG INJ 1L	8/31/27	US	J5C802	4/7/25-4/17/25
					J5C917	4/10/25-4/30/25
					J5C918	4/9/25-4/30/25
L8000	0264-7800-00	NACL INJ 0.9% 1000ML	8/31/27	US	J5C919	4/1/25-4/28/25