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.

Baxter Healthcare Corporation is issuing an Urgent Drug Recall to the user level for Bendamustine Hydrochloride Injection for the lot numbers listed below, as labels on the individual vials may become partially or fully detached from the vial while it is inside the carton. Baxter will be improving the material of the vial label to mitigate the recurrence of this issue for subsequently released lots. The product code and lot numbers affected by this issue are listed below and were distributed between 2/15/2023 and 8/3/2023 in the United States. There have been no reports of adverse events related to this issue. No hazardous situations or adverse events are expected as a result of this issue.

Product Code	Product Description	Lot Number	Expiration Date MM/DD/YYYY	NDC Number
2395B0105	Bendamustine Hydrochloride Injection 100mg/4mL	3A004A	12/31/2024	10019-079-01
		3A004B	12/31/2024	
		3B005A	01/31/2025	