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 three (3) lots of 0.9% Sodium Chloride Injection USP Product due to the potential for orange/brown particulate matter in solution or solution discoloration.

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

BBMI has identified through multiple complaints the potential for the product to contain orange/brown particulate matter in solution or solution discoloration. In the complaint investigation the particles were identified as iron oxide.

To date, there have been no reports of injury associated with the issue. If the orange/brown particulate matter or solution discoloration is observed before use, a minor delay could occur while obtaining a replacement product. If the orange/brown particulate matter or solution discoloration is not observed before use and the container is used on a patient, there is a potential for the particulate to be infused into the circulatory system. This could lead to patient harm that may require additional medical intervention and/or lead to permanent impairment or death.

NDC NUMBER	PRODUCT DESCRIPTION	LOT NUMBER	DISTRIBUTION RANGE	EXPIRATION DATE
0264-1800-32	0.9% Sodium Chloride Injection USP	J6B435	6/29/26 – 8/3/26	4/30/27
0264-1800-32	0.9% Sodium Chloride Injection USP	J6B444	2/27/26 – 7/23/26	4/30/27
0264-1800-32	0.9% Sodium Chloride Injection USP	J6B457	3/13/26 – 7/20/26	4/30/27