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.

SCA Pharmaceuticals (SCA) is announcing a recall of twelve (12) lots of compounded Heparin Sodium product due to the use of a raw material that was inadequately evaluated before use. As a result, the finished product Heparin Sodium labelling does not include the correct inactive ingredients.

This recall is being carried out at the hospital level with the knowledge of the Food and Drug Administration.

Product Description	**NDC Number**	**Lot Number**	**Beyond Use Date**	**Quantity Shipped**	**Dates Distributed**
Heparin Sodium 10 units/mL in 0.9% Sodium Chloride 500 mL Bag (5,000 units/500 mL)	70004-0650-44	1220018890	8/5/2020	373	6/3/2020 – 6/8/2020
		1220019240	8/14/2020	369	6/16/2020 – 6/23/2020
		1220019289	8/21/2020	366	6/23/2020 – 6/30/2020
Heparin Sodium 5 units/mL in 0.9% Sodium Chloride 500 mL Bag (2,500 units/500 mL)	70004-0655-44	1220018838	7/31/2020	356	6/1/2020 – 6/3/2020
		1220019244	8/17/2020	362	6/16/2020 – 6/22/2020
		1220019269	8/21/2020	369	6/22/2020 – 6/24/2020
		1220019278	8/21/2020	366	6/30/2020 – 7/1/2020
		1220019386	8/25/2020	362	6/24/2020 – 7/1/2020
Heparin Sodium 10 units/mL in 0.9% Sodium Chloride	70004-0652-46	1220018906	8/6/2020	388	6/5/2020 – 6/17/2020
		1220019055	8/11/2020	389	6/16/2020 – 6/24/2020
		1220019079	8/17/2020	388	6/24/2020 – 7/2/2020

Product Description	NDC Number	Lot Number	Beyond Use Date	Quantity Shipped	Dates Distributed
1,000 mL Bag (10,000 units/1,000 mL)		1220019457	8/24/2020	362	7/2/2020 – 7/15/2020