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.

ICU Medical Inc. is issuing a voluntary recall for one lot of Hospira intravenous (IV) solution. The affected product lot was manufactured for ICU Medical by Hospira, a Pfizer company, in July 2019 and distributed in the United States between September 2019 and October 2019. The affected product lot is listed below:

NDC Number	**Product Description**	**Lot Number***	**Expiration Date**	**Configuration**
0409-7953-09	Lactated Ringer's Injection, USP	07-514-FW	01-Jul-2021	1000 ml Flexible Container

* Note: The lot number on the shipping carton label may be followed by additional digits (Ex. 07-514-FW-XX)