From: Board of Pharmacy <pharmacy.subscriberlist@DCA.CA.GOV>
Sent: Friday, June 23, 2017 2:37 PM
To: PHARM-GENERAL@DCALISTS.CA.GOV
Cc: Pharmacy_Subscriberlist@DCA
Subject: CORRECTED Recall notice - Hospira Inc.

A notice emailed to subscribers on June 19, 2017, regarding products recalled by Hospira Inc. included two incorrect lot numbers for potassium phosphates injection.

The correct list of all products recalled by Hospira was emailed in a notice to subscribers on June 20, 2017, and is reprinted below.

Hospira, Inc., a Pfizer company,is recalling the products below due to a potential lack of sterility assurance. Microbial growth was detected during a routine simulation of the manufacturing process,which represents the potential for introduction of microorganisms into the products. The possibility of a breach in sterility assurance in distributed product, while remote,cannot be eliminated. Pfizer completed a Health Hazard Assessment, which concluded that in the event that impacted product is administered to a patient, there is a reasonable probability that the patient may experience adverse events ranging from fever, chills and malaise to severe adverse events including systemic invasive mycoses or systemic bacterial sepsis. The potential risk to a patient exposed to potentially impacted product arising from this issue is considered to be medium to high.

Please immediately check your inventory, quarantine, discontinue distribution of the affected product, and contact your wholesaler for directions.

8.4% Sodium Bicarbonate Injection, USP

NDC	Lot Numbers	Expiration Date	Strength	Configuration/Count
0409-6625-02	72109EV	12/01/2018	8.40% 50ml	1X25 VL
	72110EV	12/01/2018	8.40% 50ml	1X25 VL
	72112EV	12/01/2018	8.40% 50ml	1X25 VL
	72113EV	12/01/2018	8.40% 50ml	1X25 VL
	72114EV	12/01/2018	8.40% 50ml	1X25 VL
	73068EV	01/01/2019	8.40% 50ml	1X25 VL
	73071EV	01/01/2019	8.40% 50ml	1X25 VL
	73072EV	01/01/2019	8.40% 50ml	1X25 VL
	73224EV	01/01/2019	8.40% 50ml	1X25 VL
	73225EV	01/01/2019	8.40% 50ml	1X25 VL
	73230EV	01/01/2019	8.40% 50ml	1X25 VL
	73231EV	01/01/2019	8.40% 50ml	1X25 VL
	73232EV	01/01/2019	8.40% 50ml	1X25 VL
	73233EV	01/01/2019	8.40% 50ml	1X25 VL
	73234EV	01/01/2019	8.40% 50ml	1X25 VL
	73235EV	01/01/2019	8.40% 50ml	1X25 VL
	73236EV	01/01/2019	8.40% 50ml	1X25 VL
	73298EV	01/01/2019	8.40% 50ml	1X25 VL
	74058EV	02/01/2019	8.40% 50ml	1X25 VL

	74104EV	02/01/2019	8.40% 50ml	1X25 VL
	74105EV	02/01/2019	8.40% 50ml	1X25 VL
	74106EV	02/01/2019	8.40% 50ml	1X25 VL
	74107EV	02/01/2019	8.40% 50ml	1X25 VL
	74197EV	02/01/2019	8.40% 50ml	1X25 VL
	74198EV	02/01/2019	8.40% 50ml	1X25 VL
	74199EV	02/01/2019	8.40% 50ml	1X25 VL
	74200EV	02/01/2019	8.40% 50ml	1X25 VL
	74201EV	02/01/2019	8.40% 50ml	1X25 VL
	75171EV	03/01/2019	8.40% 50ml	1X25 VL
	75172EV	03/01/2019	8.40% 50ml	1X25 VL
	75173EV	03/01/2019	8.40% 50ml	1X25 VL
	75174EV	03/01/2019	8.40% 50ml	1X25 VL
	75175EV	03/01/2019	8.40% 50ml	1X25 VL
	75176EV	03/01/2019	8.40% 50ml	1X25 VL
	75177EV	03/01/2019	8.40% 50ml	1X25 VL

Neut™ Sodium Bicarbonate 4% additive solution

NDC	Lot Numbers	Expiration Date	Strength	Configuration/Count
0409-6609-25	75386EV	03/1/2019	4%/5ML	1X25 FTV NOVA PLUS
0409-6609-02	72226EV	12/01/2018	4%/5ML	1X25 FTV
	72236EV	12/01/2018	4%/5ML	1X25 FTV
	75382EV	03/01/2019	4%/5ML	1X25 FTV
	75383EV	03/01/2019	4%/5ML	1X25 FTV

Quelicin® Succinycholine Chloride Injection, USP

NDC	Lot Numbers	Expiration Date	Strength	Configuration/Count
0409-6629-02	74393EV	05/01/2018	200mg/10ml	1X25 FTV
	75157EV	06/01/2018	200mg/10ml	1X25 FTV
	75367EV	06/01/2018	200mg/10ml	1X25 FTV
0409-6629-25	75158EV	06/01/2018	200mg/10ml	1X25 FTV NOVAPLUS

Potassium Phosphates Injection, USP

NDC	Lot Numbers	Expiration Date	Strength	Configuration/Count
0409-7295-01	74119EV	02/01/2019	45mM	25X15ML
	74120EV	02/01/2019	45mM	25X15ML
	74121EV	02/01/2019	45mM	25X15ML
	74307EV	02/01/2019	45mM	25X15ML
	75326EV	. 03/01/2019	45mM	25X15ML
	75327EV	03/01/2019	45mM	25X15ML
	75215EV	03/01/2019	45mM	25X15ML