From: General Board of Pharmacy Subscriber List on behalf of Board of Pharmacy
To: PHARM-GENERAL@DCALISTS.CA.GOV
Subject: Recall notice - Hospira, Inc.
Date: Wednesday, March 09, 2016 4:00:53 PM

Hospira, Inc., a Pfizer company, is recalling the products identified below due to the potential contact of the products with an aromatic hydrocarbon resin.

The impacted lots were distributed January 2015 through July 2015.

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

Product	***NDC Number***	***Lot****	***Expiration Date***
70% Dextrose Injection, Partial Fill, 500mL/1000mL	0409-7918-19	48-010-JT	1DEC2016
50% Dextrose Injection, Partial Fill, 500mL/1000mL	0409-7936-19	52-002-JT	1APR2017
20% Dextrose Injection, Partial Fill, 500mL/1000mL	0409-7935-19	52-114-JT	1OCT2016
Aminosyn®-PF 7%, Sulfite-Free, An Amino Acid Injection – Pediatric formula, 500 mL	0409-4178-03	49-047-JT	1JUL2016
6% Hetastarch in 0.9% Sodium Chloride Injection, 500 mL, NOVAPLUS	0409-7248-49	50-087-JT	1FEB2017
HEXTEND®, 6% Hetastarch in Lactated Electrolyte Injection, 500mL	0409-1555-54	51-080-JT	1MAR2017
20% Mannitol I.V., Mannitol Injection, USP (20 g/100 mL), 250 mL	0409-7715-02	52-031-JT	1OCT2016
Potassium Chloride Injection (400 mEq/L), 20 mEq, 50 mL	0409-7077-14	53-006-JT	1NOV2016

*Note: The lot number may be followed by numbers from 01 to 99

---------- To unsubscribe from this email list please click on the link below and follow the instructions on the web page.

https://www.dca.ca.gov/webapps/pharmacy/subscribe.php