From: General Board of Pharmacy Subscriber List <PHARM-GENERAL@DCALISTS.CA.GOV> on behalf of Board of Pharmacy <pharmacy.subscriberlist@DCA.CA.GOV>
Sent: Friday, May 19, 2017 2:21 PM
To: PHARM-GENERAL@DCALISTS.CA.GOV
Subject: Product Recall - Hospira

Hospira, Inc., a Pfizer company, is recalling the lots below due to the potential for a foreign stopper, which is not used by Hospira McPherson site, to have been used during filling of vials. This recall is to the pharmacy level. Affected product shipped between January and February 2017.

Description	Lot #	Exp Date	NDC	Configuration
Levophed Norepinephrine Bitartrate Injection, USP 4mg/4mL (1mg per ml)	720503A	1Jun2018	0409-3375-04	10 Vials/Carton 18 cartons/case

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