From: Board of Pharmacy <pharmacy.subscriberlist@DCA.CA.GOV>
Sent: Tuesday, October 10, 2017 2:14 PM
To: PHARM-GENERAL@DCALISTS.CA.GOV
Cc: Pharmacy_Subscriberlist@DCA
Subject: Recall notice - AvKARE

AvKARE is initiating a Class III Recall on Duloxetine DR Capsule 20mg 50ct and a Market Withdrawal on Duloxetine DR Capsule 30mg 30ct and Duloxetine DR Capsule 60mg 30ct. The Recall is to the pharmacy level. This recall has been initiated due to an "Out of Spec" in stability testing with slightly elevated levels of phthalic acid. There is no indication this product is a Health Hazard to the user.

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

Product Description	NDC	Package Date	Lot & Expiration
Recalled			
Duloxetine Delayed-release Capsules 20mg 50ct	50268-283-15	06/27/2017	Lot # 18103 Exp. 11/2018
Market Withdrawal			
Duloxetine Delayed-release Capsules 30mg 30ct	50268-284-13	06/27/2017 09/26/2017	Lot # 18104 Exp. 11/2018 Lot # 18969 Exp. 02/2019
Duloxetine Delayed-release Capsules 60mg 30ct	50268-285-13	06/27/2017	Lot # 18105 Exp. 11/2018