

## Pharmacy\_Subscriberlist@DCA

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**From:** Board of Pharmacy <pharmacy.subscriberlist@DCA.CA.GOV>  
**Sent:** Tuesday, January 24, 2017 3:54 PM  
**To:** PHARM-GENERAL@DCALISTS.CA.GOV  
**Cc:** Pharmacy\_Subscriberlist@DCA  
**Subject:** Recall notice - Aurobindo Pharma USA

Aurobindo Pharma USA is recalling one lot of Venlafaxine Extended Release Capsules (37.5mg) in the United States. This recall is being carried out to the Retail level. The reason for this recall is some of the bottles were found to contain clumped/melted capsules. This lot commenced shipping to customers from August 2, 2016.

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

Hydrochloride Extended Release Capsules 37.5mg batches.