Pharmacy_Subscriberlist@DCA

From: Board of Pharmacy <pharmacy.subscriberlist@DCA.CA.GOV>
Sent: Friday, February 12, 2016 1:18 PM
To: PHARM-GENERAL@DCALISTS.CA.GOV
Cc: Pharmacy_Subscriberlist@DCA
Subject: Recall notice - Sun Pharmaceutical Industries, Ltd.

This notice is to inform you of a drug product recall involving:

Drug Product Name: Alendronate Sodium Tablets USP, 70 mg

Manufacturer: Sun Pharmaceutical Industries, Ltd.

NDC Number: 47335-638-68 (Blister of 4 Tablets)

Dosage Form: Tablets

Intended Use/Indications: Indicated for treatment and prevention of osteoporosis in postmenopausal women.

Package Type and Number of Doses/Sizes: 4 tablets packed in aluminium blister.

Product name	Batch No.	Mfg. Date	Exp. Date
Alendronate Sodium Tablets USP, 70 mg	JKP2234A	05/2015	04/2017
	JKP2235A	05/2015	04/2017

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