California Drug Recall Information

Recall Name
Pfizer Recalls **Pfizer's Effexor XR 150mg and Greeenstone's Venlafaxine HCl 150mg** **Extended-Release Capsules** **Due to Possible Presence of Tikosyn® Capsules**

Recall Date	Product Description	Recalling Firm	Recall Reason
03/06/14	**Effexor XR** (venlafaxine HCl) 150mg Extended Release Capsules • 30 ct. Caps.: **NDC 0008-0836-21** • 90 ct. Caps.: **NDC 0008-0836-22** <u>*Greenstone*-branded</u> Venlafaxine HCl Extended Release Capsules • 90 ct. Caps: **NDC 59762-0182-2**	**Pfizer, Inc.** New York, NY	*Report that one bottle of Pfizer's Effexor XR incorrectly contained one capsule of Tikosyn (dofetilide) 0.25mg.*

Recall Class	Product Identification	Distribution	Affected Dates
N/A	<u>Effexor XR</u> Lot Codes: **V130142**, exp 10/15 **V130140**, exp 10/15 <u>Greenstone</u> Lot Code: **V130014**, exp 08/15 Photos of Product Labels	**CA**, nationwide	Distribution dates: • <u>Effexor XR</u>: 3/2013 to 2/2014 • <u>Greenstone</u>: 5/2013 to 8/2013

FOR ADDITIONAL INFORMATION, PLEASE VISIT:
http://www.fda.gov/Safety/Recalls/ucm388329.htm