

Administrator Alert

Policy updates & rule clarifications for Community Based Care Facilities

Office of Licensing and Regulatory Oversight

3/12/2014

Medication Recall

Pfizer Recalls Some Antidepressants After Drug Mixup

Two lots of Pfizer's antidepressant drug Effexor XR (venlafaxine HCl) are being recalled because they may contain capsules of another drug called Tikosyn (dofetilide), which is used to treat heart rhythm disorders.

The recall also includes one lot of generic Greenstone brand venlafaxine HCl capsules, the **U.S. Food and Drug Administration** said Friday.

The agency said that unknowingly taking Tikosyn could have serious and potentially fatal consequences.

The recall is for:

- one lot of 30-count Effexor XR 150-milligram extended-release capsules,
- one lot of 90-count Effexor XR 150-milligram extended-release capsules,
- and one lot of 90-count Greenstone venlafaxine HCl 150-milligram extended-release capsules.

The Pfizer drug lot numbers are V130142 and V130140, and have an expiration date of October 2015. The Greenstone lot number is V130014, which has an expiration date of August 2015.

The voluntary recall comes after a pharmacist found that a bottle of Effexor XR contained one capsule of Tikosyn 0.25 milligram, the FDA said.

The agency advises pharmacists to immediately halt sales of the recalled lots of the drugs and notify customers who've been sold the drugs. Patients with the recalled medicines should contact their doctor and/or return them to their pharmacy.

The link to the recall announcement is listed below.

Important Drug Recall Message: http://www.drugs.com/news/pfizer-recalls-some-antidepressants-after-mixup-50690.html?utm_source=facebook&utm_medium=micro-blog&utm_campaign=DrugscomFB