



Zelnorm Recall Summary

On March 30, 2007, the FDA removed Zelnorm from the market. The Windsor Pharmacy Department determined the members that had filled a Zelnorm prescription within the previous 120 days. On April 04, 2007, the Windsor Pharmacy Department mailed a letter to both the member and their prescribing physician detailing the recall.

Date Letter Mailed	# MAPD Letters	# PDP Letters
04/06/2007	23	24

See Letter below:



Windsor Medicare Extra
7100 Commerce Way Suite 285
Brentwood, Tennessee 37027

April 1, 2007

Dear Member,

This letter is to inform you that the U.S. Food and Drug Administration (FDA) has removed Zelnorm from the market as of March 30, 2007. Our records indicate that you have filled a prescription for Zelnorm within the last 120 days.

The FDA has made this decision based on newly available information of an increased chance of heart attack, stroke and worsening heart chest pains with use of the Zelnorm compared to those treated with a sugar pill. If you are experiencing severe chest pain, shortness of breath, dizziness, sudden onset of weakness or difficulty walking or talking or other symptoms of a heart attack or stroke, contact your physician immediately.

We are also sending a notification to your prescribing physician to be sure they are aware of the Zelnorm recall. Please consult with your physician to discuss appropriate alternative treatments in regards to the Zelnorm recall.

Thank you,

Windsor Medicare Extra
Pharmacy Department

CC: MD