



**IMPORTANT
RECALL
INFORMATION**

Wellbutrin XL® Recall Summary

On March 6, 2008 GlaxoSmithKline removed Wellbutrin XL® 300mg tablets (LOT # P08A011) from the market. The Windsor Pharmacy Department determined the members that had filled a prescription for this particular NDC within the last 120 days. On March 31, 2008 the Windsor Pharmacy Department mailed a letter to the member detailing the recall.

Date Letter Mailed	# MAPD Letters	# PDP Letters
03/31/2008	4	5

See Letter below:



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March 31, 2008

Dear Member,

This letter is to inform you that GlaxoSmithKline is voluntarily recalling Wellbutrin XL® 300mg Tablets due to the possibility of embedded foreign material within the tablets.

Our records show you may have used this product in the past 120 days. If so, please discontinue using this product and return it to the manufacturer as listed below.

The chances that the use of this product posed a health risk are remote. However, as our main concern is always the health of those we serve, we advise you take the proper steps to return this product.

Product Description: Wellbutrin XL® 300mg Tablets

NDC Number: 0173-0731-01

Product Lot Numbers: P08A011

Please contact Stericycle at 1-866-324-3735 for recall instructions and further information.

As always, our Member Service department is here to help you at 1-800-316-2273 or 615-782-7878, should you have any questions.

Thank you,

Windsor Medicare Extra
Pharmacy Department