Recall Notice

VOLUNTARY DRUG RECALL

Product:	DIN:	Lot(s):
ACT OXYCODONE CR 5 mg tablets	02394170	K51225
ACT OXYCODONE CR 10 mg tablets	02394189	K51224
ACT OXYCODONE CR 20 mg tablets	02394197	K51227
ACT OXYCODONE CR 40 mg tablets	02394200	K51204
ACT OXYCODONE CR 80 mg tablets	02394219	K51228

Distributed by:
Teva Canada Limited, Toronto, Canada

July 12, 2019

Dear Pharmacist(s):

Teva Canada Limited is voluntarily recalling **ACT OXYCODONE CR 5 mg, 10 mg, 20 mg, 40 mg and 80 mg tablets**, with the lot numbers noted above.

The voluntary recall is occurring because the product lots were labeled with an incorrect expiry date.

Teva Canada is proceeding with a **Type III** recall of the lot number noted above to the **Retail Level** with the knowledge of Health Canada.

Please examine your inventory and determine if you have any remaining stock with the lot numbers noted above. If you distributed product to another wholesaler, distribution center, retailer, please notify them to cease distribution.

If you ordered product direct from Teva Canada, you will be provided a recall package that includes a returns form. **Please return all products to your point of purchase.** Your account will receive a credit for all returns. If you have any specific questions regarding the recall process, please contact Teva Canada Customer Care at 1-800-268-4129.

Teva Canada would like to thank you for your understanding and continued support.

Teva Canada Limited