

June 16, 2010

Urgent Product Recall Notification

We are requesting the return of lot # **08330-026** manufactured **Apotex** by as follows:

Level of Recall:	Retail level
Product:	Omeprazole DR 20mg
Reason:	This recall is being initiated due to a potential GMP (Good Manufacturing Practice) violation. No significant adverse health consequences are expected with these lots.
Lot #s	08330-026
Mfg	Apotex
Mfgrs. NDC #	60505-0065-00

This voluntary recall is extended to the retail level and is being made with the knowledge of the Food and Drug Administration.

This recall is only to your clinic level. You do not need to contact any clients to whom the product was dispensed.

Please fill out this sheet and fax it back to me, even if you do not have any in stock. Thanks

Our records show that product with the referenced Lot #s has been shipped to your clinic. Please examine your inventory to verify if you have product from these Lots remaining in stock. If so, **please discontinue dispensing it and promptly call Pharmedix at 1-800-486-1811, Ext. 11** to obtain the approved paperwork to return all concerned product to us for credit to your account.

Please check the appropriate box below, sign where indicated **and fax this form back to Pharmedix at 1-800-783-2038 even if you do not have any product from the indicated Lot # remaining in stock.**

Sincerely,

Tonya Tasneem

Client Services
800-486-1811x11

Clinic Code	Order Code	PMX Lot #	Rx #	Quantity

Check appropriate box:

Received affected Lot Number(s) – Do not have any stock on hand.

Received affected Lot Number(s) – Have ____ units remaining in stock.

Person verifying stock status (print): _____

Date: _____

Signature: _____