

March 2011 Recall

Urgent Product Recall Notification

We are requesting the return of lot #(s) 102209-P1, 120209-P1, 010410-P4, and 030210-P1 manufactured by **Apotex** as follows:

Level of Recall:	Voluntarily Retail level
Product:	Amlodipine Besylate 5mg
Reason:	Recalled by the manufacture Apotex to the retail level due to a routine product stability testing that identified the level of related compound-RC4 (product degradation) that exceeds the specification that may affect the product performance over shelf life. Please check your current stock of Amlodipine 5mg. You may have received one of the 4 affected lot # (s) as follows.
Lot #s	102209-P1, 120209-P1, 010410-P4 & 030210-P1
Mfg	Apotex
Mfgs. NDC #	60505-0194-03

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

Our records show that product with the referenced Lot # (s) has been shipped to your clinic in the past. Please examine your inventory to verify if you have product from these lots remaining in stock. If so, **please discontinue dispensing Amlodipine, quarantine the product 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: _____