

Urgent Product Recall Notification (Initiated 09-15-2011)

Pharmedix is requesting the return of the following product manufactured by **Jant Pharmacal Corporation** as follows:

Level of Recall:	RETAIL
Product:	Accutest Integrated Strept A Rapid Test Device (catalog# ID440)
Reason:	May have an elevated incidence of false positive results which could lead to the administration of unnecessary antibiotics
Manufacturer Lot #s	All unexpired lot numbers including STA0040091, STA0070029, STA0070075, STA0090004, 11164, 11641
Pharmedix Lot#s	All unexpired lot numbers including 10217-022, 10299-037, 10321-056, 11048-034, 11143-017, 11174-013
Manufacturer:	Jant Pharmacal Corporation
NDC Code#	63924-440-22

This voluntary recall is extended to the **RETAIL** level and is being made with the knowledge of the Food and Drug Administration (FDA). This recall is only to your **CLINIC LEVEL**. **You DO NOT need to contact any clients for whom the product was use for.**

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,

Richard J Mangini

Richard J. Mangini, PharmD
Client Services

Clinic Code	Order Code	PMX Lot #	Original Mfg. Lot #

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: _____

