



U.S. Pharmaceuticals
Pfizer Inc
235 East 42nd Street
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Pfizer Global Pharmaceuticals

Tom McPhillips
Vice President, U.S. Trade Group

URGENT: DRUG RECALL

July 8, 2010

Arthrotec[®] 75mg/200mcg (diclofenac sodium/misoprostol) Tablets

and

Arthrotec[®] 50mg/200mcg (diclofenac sodium/misoprostol) Tablets

NDC	Lot	Expiration	Strength	Configuration/Count
0025-1421-60	C090972	Feb 2013	75mg/200mcg	Bottles of 60
0025-1421-60	C090149	Sep 2012	75mg/200mcg	Bottles of 60
0025-1411-90	C080071	Jul 2011	50mg/200mcg	Bottles of 90

Dear Customer:

Pfizer Inc is recalling the above referenced lots of **Arthrotec[®] 50mg/200mcg and 75mg/200mcg Tablets**. Pfizer Inc voluntarily initiated this recall when it was determined that these lots may contain broken tablets. Please note the probability for serious adverse health consequences is low, however, if taken, could present a temporary or reversible medical hazard to the patient.

FEDERAL REGULATIONS REQUIRE THAT YOU RESPOND TO THIS RECALL, EVEN IF YOU DO NOT HAVE THE RECALLED PRODUCT. TO RESPOND, COMPLETE THE REQUESTED INFORMATION ON THE ENCLOSED POSTAGE-PAID, BUSINESS REPLY CARD AND RETURN IT TO US WITHIN FIVE (5) BUSINESS DAYS.

The recall of **Arthrotec[®] 50mg/200mcg and 75mg/200mcg Tablets** is being conducted to the **retail level**.

Our records indicate that you may have received shipment of the affected lots between **Feb 2008** and **Jan 2010**. Please check your stock immediately against the table above. If you have any of the affected product in your inventory, please stop distribution and promptly return it to Stericycle Inc.; 2670 Executive Drive; Suite A, Indianapolis, IN 46241 using the enclosed pre-paid UPS label. If you require additional shipping labels or have questions regarding the return procedure, please contact Stericycle Inc. at 1-800-805-3093.

If you have further distributed these lots to other wholesale or retail level accounts, please conduct a sub-recall to those accounts and communicate this recall information immediately. Please request that they immediately cease distribution of the affected lots and promptly return the product directly to the above address (your accounts do not need to fill out a business reply card; however, if they have inventory of the affected product, they can contact Stericycle Inc. at 1-800-805-3093 to obtain pre-paid shipping labels for product return). Further authorization is not required for product return. Reimbursement for the returned product will be made by credit memorandum. If you have any questions regarding the reimbursement, please contact your Pfizer Customer Service Representative at 1-800-533-4535.

This recall is being conducted with the knowledge of the Food and Drug Administration. We appreciate your immediate attention and cooperation and sincerely regret any inconvenience you may have been caused by this action. If you have any medical information questions regarding the product, please contact Pfizer Medical Information at 1-800-438-1985.

Sincerely,