

To: DC Managers
From: Cynthia Gaw
Date: November 3, 2010
RC: 10-126A

URGENT!!! DRUG RECALL!!! URGENT!!!

UPDATE: ADDED LOT NUMBERS

FDA/SUPPLIER CLASS OF RECALL: Not Yet Classified
LEVEL OF NOTIFICATION: McKesson Customer
SUPPLIER: Pfizer # 23386

Product Description	Lot #	Exp Date	NDC	UPC	Econo #
ARTHROTEC TAB 50/200MCG 90	C080071	07/2011	00025141190	30025141190	2130821
ARTHROTEC TAB 75/200MCG 60	C090149	09/2012	00025142160	30025142160	2130656
	C090972	02/2013			
	C091403	6/2013			
	C091600	7/2013			

Pfizer Inc is recalling the above lots because it was determined that these lots may contain broken tablets. The probability for serious adverse health consequences is low, however, if taken, could present a temporary or reversible medical hazard to the patient. This recall is to the McKesson Customer level. Product started shipping December 2009.

Please check your inventory for the above lots. Complete and return the Business Reply Card within five business days even if you do not have product in stock. If you have any of the affected products in your inventory, please stop distribution and promptly return to the address below use the pre-paid UPS label. If you require additional shipping labels, need a Business Reply Card or have questions regarding the return procedure, please contact Stericycle Inc. at 1-800-805-3093.

Stericycle Inc.
2670 Executive Drive
Suite A
Indianapolis, IN 46241

McKesson customers are to check their inventory and return to Stericycle.

If you have any questions regarding the reimbursement, please contact your Pfizer Customer Service Representative at 1-800-533-4535. If you have questions about this recall, contact Stericycle at 800-805-3093.

This recall is being conducted with the knowledge of the FDA.

Information contained in this document was provided by Pfizer.

FOR McKESSON USE ONLY:

- Please follow Recall Guidelines as outlined in the Reclamation SOP for a CLASS II RECALL.
- PROCESS TO THE RETAIL LEVEL.