McNeil Consumer Healthcare Over-The-Counter Products: Recall(2)

U.S. Food & Drug Administration

[UPDATED 06/17/2010] Additional product lots of Benadryl Allergy Ultratab Tablets and Extra Strength Tylenol Rapid Release Gels added to the January 15 2010 Recall.

McNeil and FDA notified healthcare professionals of an expansion of the December 2009 recall. McNeil Consumer Healthcare has now applied broader criteria to identify and remove all product lots that it believes may have the potential to be affected, even if they have not been the subject of consumer complaints. Consumers who purchased product from the lots included in this recall should stop using the product and contact McNeil Consumer Healthcare for instructions on a refund or replacement. The affected product names and lot numbers for the recalled products can be found in firm's Press Release. Any adverse reactions may also be reported to the FDAâ€™s MedWatch Program by fax at 1-800-FDA-0178, by mail at MedWatch, FDA, 5600 Fishers Lane, Rockville, MD 20852-9787, or on the MedWatch website at www.fda.gov/medwatch [1].

[06/15/2010 - Press Release [2] - McNeil Consumer Healthcare]
[01/15/2010 - Press Release [3] - McNeil Consumer Healthcare]
[12/18/2009 - Press Release [4] - McNeil Consumer Healthcare]

Previous Medwatch alert:

[Posted 12/28/2009 - Tylenol Arthritis Pain Caplet, 100 count bottles: Recall of all lots [5]]

SOURCE [6]

Source URL (retrieved on *12/20/2013 - 4:25pm*):
http://www.mdtmag.com/news/2010/06/mcneil-consumer-healthcare-over-counter-products-recall2

Links:
[1] http://www.fda.gov/medwatch
[2] http://www.fda.gov/Safety/Recalls/ucm215921.htm
[3] http://www.fda.gov/Safety/Recalls/ucm197746.htm
[4] http://www.fda.gov/Safety/Recalls/ArchiveRecalls/2009/ucm195690.htm
[5] http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm195704.htm
[6] http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm197740.htm