



June 15, 2010

Dear Healthcare Professional:

I am writing to inform you that McNeil Consumer Healthcare, Division of McNEIL-PPC, Inc. is voluntarily recalling five product lots as an addition to the list of products included in the company's January 15th, 2010 product recall. The additional lots involved are four product lots of BENADRYL® ALLERGY ULTRATAB™ TABLETS, 100 count, sold in the U.S.; and one product lot of EXTRA STRENGTH TYLENOL® Rapid Release Gels, 50 count, sold in the U.S., Trinidad and Tobago, Bermuda, and Puerto Rico (FULL RECALLED PRODUCT LIST BELOW). This recall is being conducted with the knowledge of the U.S. Food and Drug Administration. McNeil is providing the following information to consumers on the [www.mcneilproductrecall.com](http://www.mcneilproductrecall.com) Web site.

**MCNEIL CONSUMER HEALTHCARE RECALLS FOUR PRODUCT LOTS OF BENADRYL® ALLERGY ULTRATAB™ TABLETS, 100 COUNT, AND ONE PRODUCT LOT OF EXTRA STRENGTH TYLENOL® RAPID RELEASE GELS, 50 COUNT**

Fort Washington, PA (June 15, 2010) – McNeil Consumer Healthcare, Division of McNEIL-PPC, Inc., is recalling five product lots as an addition to the list of products included in the company's January 15th, 2010 product recall. The additional lots involved are four product lots of BENADRYL® ALLERGY ULTRATAB™ TABLETS, 100 count, sold in the U.S.; and one product lot of EXTRA STRENGTH TYLENOL® Rapid Release Gels, 50 count, sold in the U.S., Trinidad and Tobago, Bermuda, and Puerto Rico (FULL RECALLED PRODUCT LIST BELOW). This recall is a follow-up to the product recall that McNeil Consumer Healthcare announced on January 15th, 2010 and is being taken because the products were inadvertently omitted from the initial recall action. McNeil Consumer Healthcare identified the omission and informed the U.S. Food and Drug Administration (FDA) of its decision to add these product lots to the recall list. All of these products were produced before the January 15th, 2010 recall. Since January, McNeil Consumer Healthcare has continued to analyze and evaluate 2,4,6-tribromoanisole (TBA) and has shared that information with the FDA. This further analysis confirms that the risk of serious adverse medical events is remote. This recall is being conducted with the knowledge of the FDA.

BENADRYL® ALLERGY ULTRATAB™ TABLETS is sold over-the-counter and is indicated for the relief of allergy associated symptoms such as sneezing, runny nose, itchy throat, and itchy and watery eyes. EXTRA STRENGTH TYLENOL® Rapid Release Gels is sold over-the-counter and is indicated for the temporary reduction of fever and for the temporary relief of minor aches and pains due to headache, muscular aches, backache, minor arthritis pain, the common cold, toothache, pre-menstrual and menstrual cramps, and flu.

The January 15th, 2010 recall was initiated as a result of consumer complaints of a musty or moldy odor that has since been linked to the presence of trace amounts of a chemical called 2,4,6-tribromoanisole (TBA). After a thorough investigation, it was determined that the source of TBA was the result of a breakdown of a chemical that is applied to wood used to build wooden pallets that transport and store product packaging materials.

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. For these instructions or information regarding how to return or dispose of the product, consumers should log on to the Internet at [www.mcneilproductrecall.com](http://www.mcneilproductrecall.com) or call 1-888-222-6036 (Monday-Friday 8 a.m. to 8 p.m. Eastern Time, and Saturday-Sunday 9 a.m. to 5 p.m. Eastern Time). Requests for reimbursement can be made at [www.mcneilproductrecall.com](http://www.mcneilproductrecall.com). Consumers who have medical concerns or questions should contact their healthcare provider.

Any adverse reactions may also be reported to the FDA's MedWatch Program by fax at 1-800-FDA-0178, by mail at MedWatch, HF-2, FDA, 5600 Fishers Lane, Rockville, MD 20852-9787, or on the MedWatch Web site at <http://www.fda.gov/safety/medwatch/default.htm>.

The product lot numbers for the recalled products can be found on the side of the bottle label.

If you have any questions yourself, please call our Medical Affairs Department at **1-866-948-6983** (available Monday-Friday 8 a.m. to 8 p.m. Eastern Time), or visit our dedicated Web site for Healthcare Professionals at [TylenolProfessional.com](http://TylenolProfessional.com).

Sincerely,



Edwin K. Kuffner, MD  
Vice President, Medical Affairs  
McNeil Consumer Healthcare

#### FULL RECALLED PRODUCT LIST:

Product Code	Product Name	Lot Number	UPC Code
<b>TYLENOL® EXTRA STRENGTH RAPID RELEASE GELCAPS</b>			
4885000	TYLENOL® EXTRA STRENGTH RAPID RELEASE GELCAPS 50 ct	ASA202	300450488503
<b>BENADRYL® ALLERGY ULTRATAB™ TABLETS</b>			
1703302	BENADRYL® ALLERGY ULTRATAB™ TABLETS 100 ct	ABA022	312547170338
		ABA264	312547170338
		ADA194	312547170338
		AJA008	312547170338

If you no longer wish to receive fax messages from this sender, please call 888-739-4920, enter extension 2283 and follow remaining voice prompts.