Insight Issues Voluntary Nationwide Recall of Ting® 1% Tolnaftate Athlete's Foot Spray Due to the Presence of Benzene

Date: 02/02/2024

About this recall:
Insight is voluntarily recalling 2 lots of Ting® 1% Tolnaftate Athlete's Foot Spray Antifungal Spray Liquid (NDC 63736819-05) to the consumer level. A recent review by the manufacturer and their third-party lab found that samples from 2 lots of the product contained elevated levels of benzene. The benzene came from the propellant that sprays the product out of the can. Benzene is not an ingredient in any Ting® Antifungal Spray products and is classified as a human carcinogen (a substance the can cause cancer). No other lots of Ting 1% Tolnaftate Athlete's Foot Spray and no other Ting Antifungal Spray Liquid products are in the scope of this recall. The recalled.

For lot recalls, the lot/batch information and expiration date for the recalled product can be viewed by clicking on the link to the FDA Recall Notification found on the following page. Lot codes for this product are located on the bottom of the can.

Ting 1% Tolnaftate Athlete's Foot Spray is an over the counter (OTC) antifungal used for (1) the treatment and prevention of most athlete's foot, and (2) it relieves itching, scaling, burning and discomfort that can accompany athlete's foot.

What this means to you:
Exposure to benzene can occur through the lungs, mouth, or skin and potentially may result in blood cancers or blood disorders which can be life threatening. Benzene is found throughout the environment, and humans have daily exposures to it both indoors and outdoors from multiple sources. These products have been recalled out of an abundance of caution.

The company will offer reimbursement for consumers who have purchased recalled Ting® 1% Tolnaftate Athlete's Foot Spray. Consumers can contact Insight Pharmaceuticals via e-mail at medicalaffairs@prestigebrands.com, through its website at https://www.prestigebrands.com/contact, or by phone at (800) 344-7239 Monday through Friday 8:30 am to 5:30 pm eastern time to receive a full refund by providing a picture of the bottom of the can of the product with a recalled lot number. **Consumers that have product which is being recalled should stop using the product immediately and appropriately discard after taking the picture.** Consumers with questions regarding this recall can also contact Insight at the email, website, or phone number listed above. Consumers should contact

their healthcare provider if they have experienced any problems that may be related to taking or using this antifungal product.

Adverse reactions or quality problems experienced with the use of this product may be reported to the FDA's MedWatch Adverse Event Reporting program either online, by regular mail, or by fax.

- Complete and submit the report Online.
- Regular Mail or Fax: Download form or call 1-800-332-1088 to request a reporting form, then complete and return to the address on the pre-addressed form, or submit by fax to 1-800-FDA-0178.

This recall is being conducted with the knowledge of the United States (US) Food and Drug Administration (FDA).

For more information regarding this FDA Recall Notification, please refer to the FDA website: https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/insight-pharmaceuticals-issuesvoluntarynationwide-recall-tingr-1-tolnaftate-athletes-foot-spray

FDA contact information for reporting adverse events/quality complaints can be reached online at https://www.accessdata.fda.gov/scripts/medwatch/index.cfm?action=reporting.home or by calling the FDA at 1888-INFO-FDA (1-888-463-6332) and then selecting prompt #2.

