Date: October 1, 2021

URGENT LOTRIMIN® AND TINACTIN® SPRAY RECALL

Dear [Retailer],

This letter is to inform you that Bayer Consumer Health is voluntarily recalling all unexpired Lotrimin® Anti-Fungal (AF) and Tinactin® spray products listed below with lot numbers beginning with TN, CV or NAA distributed between September 2018 to September 2021. The recall is due to low levels of residual solvent benzene detected in some samples of the final product. Benzene is not an ingredient in any of Bayer Consumer Health products. It is important to note that Bayer's decision to voluntarily recall these products is a precautionary measure and that the levels detected are not expected to cause adverse health consequences in consumers.

The affected Lotrimin® and Tinactin® spray products are over-the-counter antifungal products, sold individually or in combo packs. The impacted products are:

PRODUCT DESCRIPTION	UPC
LOTRIMIN AF JI SPRAY POWDER 133G 36PK	00311017410318
LOTRIMIN AF POWDER SPRAY 133G 36PK	00311017410257
LOTRIMIN AF DEO POWDER SPRAY 133G 36PC	00311017410233
LOTRIMIN LIQUID SPRAY 133G (4.6OZ)	00041100407887
LOTRIMIN DAILY PREV DEO POWDER SPRAY 160G	00041100587206
LOTRIMIN DAILY PREV DEO POWDER SPRAY 133g	00041100590367
LOTRIMIN AF POWDER SPRAY 133G (3 Pack)	00041100585943
LOTRIMIN AF LIQUID SPRAY 133G (3 Pack)	00041100585936
LOTRIMIN DAILY PREV DEO POWDER SPRAY (3 Pack)	00041100589613
LOTRIMIN PREV SPRAY+ULTRA AF CRM (ECOM Pack)	00041100590756
TINACTIN JI SPRAY 133G	00311017410073
TINACTIN AF POWDER SPRAY 133G	00311017410097
TINACTIN AF LIQUID SPRAY 150G (CLD)	00311017410059
TINACTIN AF DEODORANT POWDER SPRAY 133G	00311017410004

There are no issues of concern with Lotrimin®/Tinactin® creams, including Lotrimin® Ultra, or any other Bayer products.

Product images and information on which lot numbers fall under this recall are available at https://livewell.bayer.com/document/2011.

Bayer asks that you take the following actions:

- Immediately examine your inventory and quarantine products subject to recall. In addition, if you may have further distributed this product, please identify your customers, and notify them at once of this product recall. Your notification to your customers may be enhanced by including a copy of this recall notification letter.
- Complete and return the enclosed response form as soon as possible via e-mail to Recall@qualanex.com or fax to 1-847-737-3719.
- When returning to Qualanex, a return authorization will be issued and e-mailed to you to be included in your return.
- Follow the instructions on the response form to arrange for return to Qualanex of any identified impacted product.
- If you receive questions from your consumers, have them contact Bayer at 1-866-360-3266.

If you have any general questions regarding the return of this product, please contact Qualanex via e-mail at Recall@qualanex.com or toll-free at 888-280-2043.

Bayer's press release regarding this announcement will be available at: https://livewell.bayer.com/document/2006.

This recall is being conducted with the knowledge of the U.S. Food and Drug Administration.

We value the relationships we have with our customers and greatly appreciate your patience through this process. If you have any additional questions, please reach out to your Bayer Sales representative.

Sincerely,

Karin Ann Payne Digitally signed by Karin Ann Payne
Date: 2021.10.01 12:58:51 -04'00'

Karin Ann Payne
Quality, Region U.S. and Canada