No Varnish
No Text Area

LOTRIMIN ULTRA
PRESCRIPTION STRENGTH

Contains the Drug: BUTENAFINE HYDROCHLORIDE

LOTRIMIN ULTRA®

BUTENAFINE HYDROCHLORIDE CREAM 1% ANTIFUNGAL

PRESCRIPTION STRENGTH

CLINICALLY PROVEN TO CURE MOST
ATHLETE'S FOOT
BETWEEN THE TOES

RELIEVES
✓ ITCHING
✓ BURNING
✓ CRACKING

1 WEEK
TREATMENT OPTION
FOR ATHLETE'S FOOT
SEE DIRECTIONS

NO MESS
APPLICATOR

NET WT 20g (0.70 OZ)

LOTRIMIN ULTRA
PRESCRIPTION STRENGTH

LOTRIMIN ULTRA®

Actual Size

LOT:
EXP:

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Contains the Drug: BUTENAFINE HYDROCHLORIDE

LOTRIMIN ULTRA
PRESCRIPTION STRENGTH

Drug Facts

Active Ingredient — **Purpose**
Butenafine hydrochloride 1% ..... Antifungal

Uses
- cures most athlete's foot between the toes. Effectiveness on the bottom or sides of feet is unknown.
- relieves itching, burning, cracking, and scaling which accompany this condition

Warnings
For external use only

Do not use
- on nails or scalp
- in or near the mouth or the eyes
- for vaginal yeast infections

When using this product do not get into the eyes. If eye contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if too much irritation occurs or irritation gets worse

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions
- adults and children 12 years and older:
  - wash the affected skin with soap and water and dry completely before applying
  - to open the applicator, twist the base to the OPEN position, squeeze tube to dispense cream.
  - for athlete's foot between the toes: apply to affected skin between and around the toes twice a day for 1 week (morning and night), or once a day for 4 weeks, or as directed by a doctor. Wear well-fitting, ventilated shoes. Change shoes and socks at least once daily.
  - wipe applicator tip with a tissue and close the applicator by twisting the base to the OFF position
  - wash hands after each use
- children under 12 years: ask a doctor

Other information
- store between 20° to 25°C (68° to 77°F)

Inactive ingredients
benzyl alcohol, cetyl alcohol, glycerin, glyceryl monostearate SE, polyoxyethylene (23) cetyl ether, propylene glycol dicaprylate, purified water, sodium benzoate, stearic acid, tolpanine, white petrolatum

Questions? 1-888-269-3356 or visit us at www.lotrimin.com

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Product of Japan
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Bayer

Recyclable in PAPER BOX MAGNAPAK TUBE

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Actual Size 100%

FDA64704-6

DRUG FACTS INFO (updated 5/18/18)		
CATEGORY	REQUIRED	ACTUAL IN FILE
Drug Facts Title	≥8 pt **	8 pt
Drug Facts Continued	8 pt	•
Headings*	8 pt+[illegible]	8 pt
Subheadings	≥8 pt	8 pt
Body Text	≥8 pt	8 pt
Bullets	5 pt	5 pt
Leading	≥0.5 pt	0.5 pt
Hairlines	0.5 pt rule	0.5 pt
Box and Underlines†	>0.5 pt rule	1 pt
Characters Per Inch	No more than 35	≤30

* Headings are to be at least 2pts larger than Body Text.
** Drug Facts Title is to be larger than all other type size used in DFP
† FDA Recommends 1pt rule

Reference ID: 5411355