LIQUID ANTIFUNGAL TREATMENT- tolnaftate liquid
Hudson Health LLC

Comfort Zone Liquid Antifungal Treatment

Active ingredient

Tolnaftate 1%

Purpose

Antifungal

Uses

for the treatment of athlete's foot (tinea pedis) and ringworm (tinea corporis)

for effective relief of itching, scaling, cracking, burning and redness.

Warnings

For external use only.

Flammable

keep away from fire or flame.

Do not use

on children under 2 years of age unless directed by a doctor.

When using this product

avoid contact with the eyes.

Stop use and consult a doctor if

- irritation occurs or if there is no improvement within 4 weeks

Keep out of reach of children.

If swallowed, get medical help, or contact a Poison Control Center right away.

Directions

- wash the affected area and dry thoroughly.
- apply a thin layer of the product over affected area twice daily (morning and night) or as directed by a doctor.
- supervise children in the use of this product.
- for athlete's foot: Pay special attention to spaces between the toes; wear well-fitting,

ventilated shoes, and change shoes and socks at least once daily.
- for athlete's foot and ringworm, use daily for 4 weeks.
- if condition persists longer, consult a doctor.
- this product is not effective on the scalp or nails.

Other information

- Store at room temperature 15°-30°C (59° - 86°F)
- Keep tightly closed when not in use

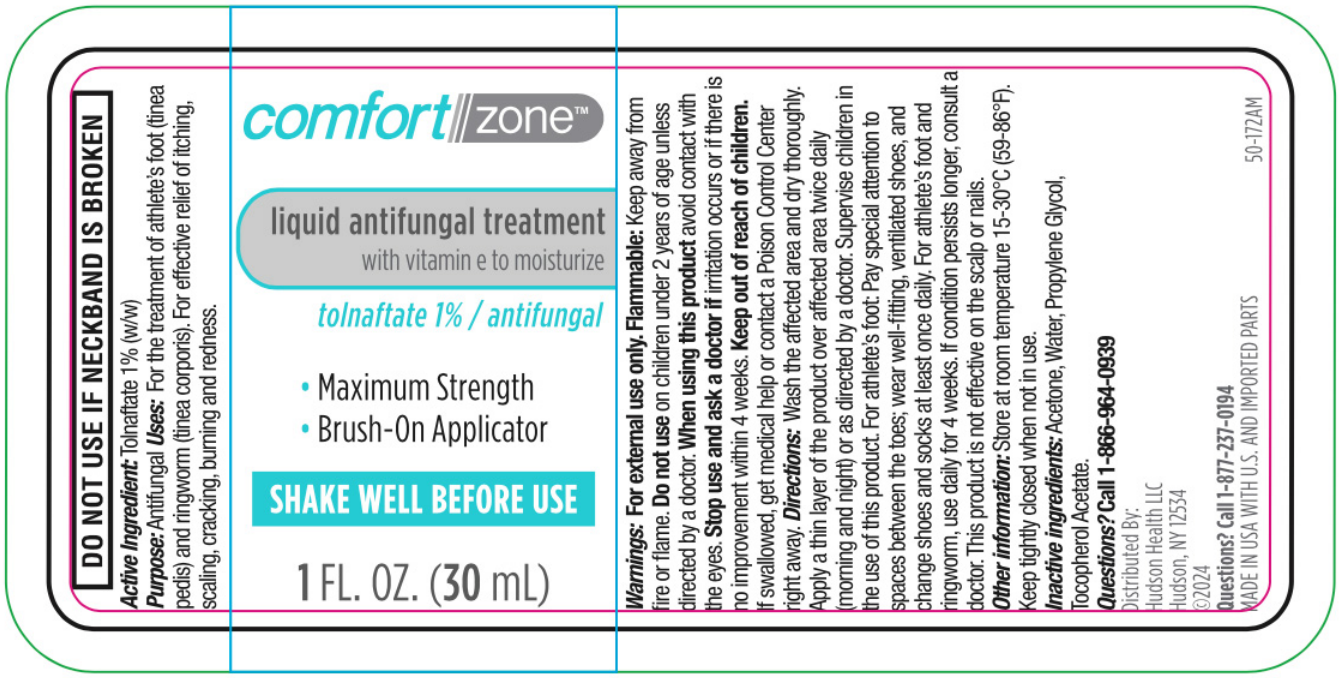
Inactive ingredient

Acetone, Water, Propylene Glycol, Tocopherol Acetate.

Questions?

Call 1-866-964-0939

Principal Display Panel



Comfort zone

liquid antifungal treatment

with vitamin e to moisturize

TOLNAFTATE 1%/ANTIFUNGAL

Maxiumum Strength

Brush- On Applicator

SHAKE WELL BEFORE USE

1 FL. OZ. (30 mL)

LIQUID ANTIFUNGAL TREATMENT

tolnaftate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72446-012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)	TOLNAFTATE	10 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ACETONE (UNII: 1364PS73AF)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72446-012-01	1 in 1 CARTON	03/29/2024	
1		30 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	03/29/2024	

Labeler - Hudson Health LLC (081276171)

Revised: 4/2024

Hudson Health LLC