LUCKY SUPER SOFT ANTIFUNGAL FOR ATHLETES FOOT- tolnaftate powder Delta Brands, Inc

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Lucky Antifng Pwdr

Active Ingredient

Tolnaftate 1%

Purpose

Antifungal

Uses

■ proven clinically effective in the treatment of most athlete's foot (tinea pedis) and ringworm (tinea corporis)
■ helps prevent most athlete's foot with daily use
■ for effective relief of itching, burning, and cracking

Warnings

For external use only

Flammable: Do not use while smoking or near heat or flame

Do not use

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

When using this product

■ avoid contact with eyes ■ use only as directed. Intentional misuse by deliberetely concentrating and inhaling contents can be harmful or fatal ■ contents under pressure. Do not puncture or incinerate.

Do not stire at a temperature above 120°F (49°C)

Stop use and ask a doctor if

■ irritation occurs ■ 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 affected area and dry thoroughly \blacksquare shake can well and spray a thin layer over affected area twice daily (morning and night) \blacksquare supervise children in the use of this product \blacksquare 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 a day \blacksquare use daily for 4 weeks, if conditions persist longer, ask a doctor \blacksquare to prevent athlete's foot, apply once or twice daily (morning and/or night) \blacksquare this product is not effective on the scalp or nails \blacksquare in case of clogging, clean nozzle with pin

Other information

 \blacksquare store between 68° to 77°F (20° to 25°C)

Inactive ingredients

alcohol denat., butane, fragrance, propane, triethanolamine, zea mays (corn) starch

Package Label



SHAKE WELL BEFORE USING

PURE CORNSTARCH
ANTIFUNGAL
POWDER
SPRAY

For Athlete's Foot

TOLNAFTATE 1%
100% TALC-FREE!

*Compare to the Active Ingredient in Tinactin®

Cures & Prevents Most Athlete's Foot Relieves Itching & Burning

NET WT. 3 OZ. (85 g)

Drug Facts

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daily (morning and night) ■ 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 ■ use daily for 4 weeks; if condition
persists longer, ask a doctor ■ to prevent athlete's foot, apply once
or twice daily (morning and/or night) ■ this product is not effective
on the scalp or nails ■ in case of clogging, clean nozzle with pin

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*This product is not manufactured or distributed by the Bayer Corporation, owner of the registered trademark Tinactin®

ITEM NO: 11386

Manufactured for DELTA BRANDS INC.
Larchmont, NY 10538 USA www.deltabrands.com

Made in China



LUCKY SUPER SOFT ANTIFUNGAL FOR ATHLETES FOOT

tolnaftate powder

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:20276-108

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TOLNAFTATE (LINII: 06KB629TKV) (TOLNAFTATE - LINII: 06KB629TKV)	ΤΟΙ ΝΑΕΤΑΤΕ	1 a in 100 a

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
BUTANE (UNII: 6LV4FOR43R)	
PROPANE (UNII: T75W9911L6)	
TROLAMINE (UNII: 903K93S3TK)	
STARCH, CORN (UNII: O8232NY3SJ)	

F	Packaging						
#	tem Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:20276-108- 85	85 g in 1 CAN; Type 0: Not a Combination Product	02/19/2018				
2	NDC:20276-108- 57	57 g in 1 CAN; Type 0: Not a Combination Product	06/01/2022				

Marketing In	Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph final	part333C	02/19/2018			

Labeler - Delta Brands, Inc (102672008)

Revised: 6/2022 Delta Brands, Inc