MEIJER MAXIMUM STRENGTH JOCK ITCH RELIEF ANTIFUNGALtolnaftate spray Formulated Solutions, LLC

Meijer Maximum Strength Jock Itch Relief Tolnaftate Antifungal

Drug Facts

Active ingredient

Tolnaftate 1%

Purpose

Antifungal

Uses

- cures most jock itch
- for effective relief of itching, chafing and burning

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 the eyes
- use only as directed intentional misuse by deliberately concentrating and inhaling contents can be harmful or fatal
- contents under pressure. Do not punture or incinerate. Do not store at temperature above 120°F (49°C).

Stop use and ask a doctor if

- irritation occurs
- there is no improvement within 2 weeks

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Directions

- wash affected area and dry thoroughly
- shake can well and spray a thin layer over affected area twice daily (morning and night)
- supervise children in the use of this product
- use daily for 2 weeks; if condition persist longer, ask a doctor
- this product is not effective on the scalp or nails
- In case of clogging, clean nozzle with a pin

Other information

store at 20-25°C (68-77°F)

Inactive Ingredients

alcohol denat, aluminum starch octenylsuccinate, isobutane, isopropyl myristate, propylene carbonate, sorbitan monooleate, stearalkonium hectorite, talc

Questions or comments?

1-800-719-9260

Package Labeling:

NDC 41250-695-90

meijer

Compare to Tinactin® **Antifungal Powder Spray**

active ingredient*

Maximum Strength

Jock Itch Relief

Tolnaftate 1% • Antifungal

Powder Spray · Goes On Dry

Cures Most Jock Itch

Relieves Itching, Burning & Chafing

NETWT 4.6 OZ (130 g)

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

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: 69590 6E F1

tolnaftate spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:23667-809

Route of Administration TOPICAL

Active Ingredient/Active Moiety

I	Ingredient Name	Basis of Strength	Strength
I	TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)	TOLNAFTATE	10 mg in 1 g

Inactive Ingredients		
Ingredient Name	Strength	
ALCOHOL (UNII: 3K9958V90M)		
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: 19PJ0O6294)		
ISOBUTANE (UNII: BXR49TP611)		
ISOPROPYL MYRISTATE (UNII: ORE8K4LNJS)		
PROPYLENE CARBONATE (UNII: 8D08K3S51E)		
SORBITAN MONOOLEATE (UNII: 06XEA2VD56)		
STEARALKONIUM HECTORITE (UNII: OLX698AH5P)		
TALC (UNII: 7SEV7J4R1U)		

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:23667-809-	130 g in 1 CAN; Type 0: Not a Combination	01/21/2017	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M005	01/21/2017		

Labeler - Formulated Solutions, LLC (143266687)

Establishment				
Name	Address	ID/FEI	Business Operations	
Formulated Solutions, LLC		143266687	manufacture(23667-809)	

Revised: 12/2023 Formulated Solutions, LLC