TOLNAFTATE- foot odor control spray aerosol, spray HEB

HEB Foot Odor Control Powder Spray

Active ingredient

Tolnaftate 1%

Purpose

Antifungal

Uses

- prevents recurrence of athlete's foot (tinea pedis) with daily use
- cures mose athlete's foot (tinea pedis) and ringworm (tinea corporis)
- relieves sypmtoms of athlete's foot, including itching, burning and cracking

Warnings

For external use only.

Flammable:

Keep away from heat, sparks and open flame. Contents under pressure. Do not puncture or incinerate. Do not store at temperature above 120°F. Intentional misuse by deliberately concentrating and inhaling contents cans be harmful or fatal.

When using this product

• avoid contact with the eyes or mouth

Stop use and ask a doctor if

- irritation occurs
- no improvement within 4 weeks

Do not use

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

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
- shake can well and spray a thin layer over affected area twice 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 conditions persist, consult a doctor
- to prevent athlete's foot, apply once or twice daily (morning and/or night)
- in case of clogging, clear nozzle under running water

Other information

store between 20° and 30°C (68°F and 86°)

Inactive ingredients

disteardimonium hectorite, fragrance, isobutane, isopropyl myristate, SD alcohol 40-B, sodium bicarbonate

Questions?

call 1-866-964-0939

Principal Display Panel

HEB

Foot Odor Control

Tolnaftate 1%

Antifungal Powder Spray

- Reduces Foot Odor
- Cures Most Athlete's Foot

NET WT 4 OZ (113g)



TOLNAFTATE

Product Information

foot odor control spray aerosol, spray

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:37808-514

Route of Administration TOPICAL

Active Ingredient/Active Moiety

SODIUM BICARBONATE (UNII: 8MDF5V39QO)

Ingredient Name

Basis of Strength

TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)

TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)

Inactive Ingredients			
Ingredient Name	Strength		
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)			
ISOBUTANE (UNII: BXR49TP611)			
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)			
ALCOHOL (UNII: 3K9958V90M)			

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
	NDC:37808-514- 40	113 g in 1 CAN; Type 0: Not a Combination Product	01/05/2012			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M005	01/05/2012			

Labeler - HEB (007924756)

Revised: 1/2024 HEB