TINACTIN- tolnaftate aerosol, powder
Bayer HealthCare LLC

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.

Tinactin ®

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 puncture or incinerate. Do not store at temperature above 120°F.

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.

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 persists longer, ask a doctor
- this product is not effective on the scalp or nails

Other information

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

Inactive ingredients

butylated hydroxytoluene, isobutane, PPG-12-buteth-16, SD alcohol 40-B (11% v/v), talc

Questions?

1-866-360-3266

PRINCIPAL DISPLAY PANEL - 133g Can Label

NDC 11523-0072-5

TOUGH ACTIN' ®

Tinactin®

ANTIFUNGAL

tolnaftate

Cures most

jock itch

POWDER SPRAY

goes on dry

Relieves itching,

burning & chafing

NET WT 133g (4.6 oz)

TINACTIN

tolnaftate aerosol, powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11523-0072
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 g

Inactive Ingredients	
Ingredient Name	**Strength**

BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
ISOBUTANE (UNII: BXR49TP611)	
PPG-12-BUTETH-16 (UNII: 58CG7042J1)	
TALC (UNII: 7SEV7J4R1U)	

Product Characteristics

Color	white (white to off-white)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11523-0072-5	133 g in 1 CAN; Type 0: Not a Combination Product	09/23/1993	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333C	09/23/1993	

Labeler - Bayer HealthCare LLC (112117283)

Revised: 9/2023 Bayer HealthCare LLC