# CANESPIES- tolnaftate cream OPMX LLC

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#### **ACTIVE INGREDIENTS**

**TOLNAFTATE 1%** 

#### **PURPOSE**

**ANTIFUNGAL** 

#### **USES**

- PROVEN CLINICALLY EFFECTIVE IN THE TREATMENT OF ATHLETE'S FOOT (TINEA PEDIS), JOCK ITCH (TINEA CRURIS), RINGWORM (TINEA CORPORIS)
- PROVEN EFFECTIVE IN THE PREVENTION OF ATHLETE'S FOOT
- EFFECTIVELY SOOTHES AND RELIEVES ITCHING ASSOCIATED WITH JOCK ITCH, SCALY SKIN BETWEEN THE TOES AND BURNING FEET

#### **WARNINGS**

For external use only.

STOP USE AND ASK A DOCTOR IF

**IRRITATION OCCURS** 

THERE IS NO IMPROVEMENT WITHIN 4 WEEKS.

DO NOT USE ON CHILDREN UNDER 2 YEARS OF AGE EXCEPT UNDER THE ADVICE AND SUPERVISION OF A DOCTOR.

WHEN USING THIS PRODUCT AVOID CONTACT WITH EYES.

### Keep out of reach of children

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

#### **DIRECTIONS**

- WASH AFFECTED AREA AND DRY THOROUGHLY
- APPLY A THIN LAYER 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, VENTILLATED SHOES AND CHANGE SHOES AND SOCKS AT LEAST ONCE DAILY
- FOR ATHLETE'S FOOT, AND RINGWORM USE DAILY FOR 4 WEEKS, FOR JOCK ITCH USE DAILY FOR 2 WEKS. IF CONDITION PERSISTS LONGER CONSULT A DOCTOR
- THIS PRODUCT IS NOT EFFECTIVE ON THE SCALP OR NAILS

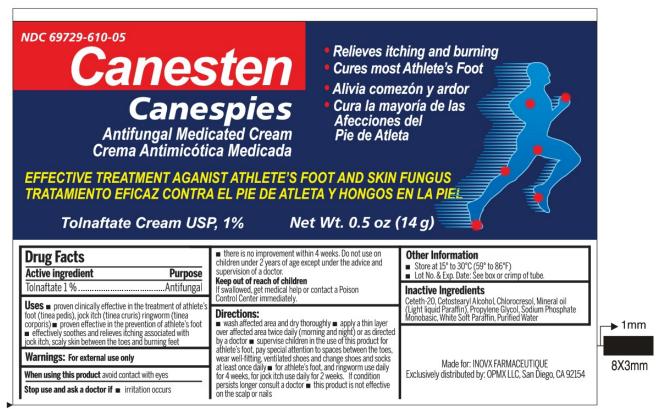
#### OTHER INFORMATION

- STORE AT 15° TO 30°C (59°F TO 86°F)
- LOT NO. & EXP. DATE: SEE BOX OR CRIMP OF TUBE

#### **INACTIVE INGREDIENTS**

CETETH-20, CETYL ALCOHOL, CHLOROCRESOL, MINERAL OIL, PROPYLENE GLYCOL, SODIUM PHOSPHATE MONOBASIC, WHITE SOFT PARAFFIN, PURIFIED WATER

NDC 69729-610-05



5 mm



## **CANESPIES**

tolnaftate cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69729-610

Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)	TOLNAFTATE	1 a in 100 a	

Inactive Ingredients		
Ingredient Name	Strength	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)		
MINERAL OIL (UNII: T5L8T28FGP)		
PARAFFIN (UNII: 1900E3H2ZE)		
CETETH-20 (UNII: 1835H2IHHX)		

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM (UNII: 3980JIH2SW)	
CHLOROCRESOL (UNII: 36W53O7109)	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69729-610- 05	1 in 1 BOX	07/26/2022	
1	NDC:69729-610- 01	14 g in 1 TUBE; 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	07/26/2022	

## **Labeler -** OPMX LLC (029918743)

Revised: 5/2024 OPMX LLC