MICONATATE- miconazole nitrate and tolnaftate PureTek Corporation

Miconatate[™]

Kit

Mycozyl AP[™], Miconazole Nitrate 2%, Antifungal Powder (3 oz / 85 g) Mycozyl AL[™], Tolnaftate 1%, Antifungal Liquid (1 fl oz / 30 mL)

Mycozyl AP

Active ingredient

Miconazole Nitrate 2%

Purpose

Antifungal

Uses

- for the treatment of most athlete's foot (tinea pedis), jock itch (tinea cruris), ringworm (tinea corporis).
- relieves itching, burning, cracking, scaling, and discomfort which accompany these conditions.

Warnings

For external use only.

Do not use on

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

When using this product

• do not get into eyes

Stop use and ask a doctor if

- for athlete's foot and ringworm irritation occurs or there is no improvement within 4 weeks
- for jock itch irritation occurs or 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

- clean the affected area and dry thoroughly
- apply a thin layer of the product 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, ventilated 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 weeks
- if condition persists longer, consult a doctor. This product is not effective on the scalp or nails.

Other information

- protect from freezing
- avoid excessive heat
- do not use if package is damaged

Inactive ingredients

Aleurites Moluccana (Kukui) Seed Oil, Aloe Barbadensis (Aloe Vera) Leaf Juice Powder, Bisabolol, Carthamus Tinctorius (Safflower) Oleosomes, Fragrance, Nylon-12, Silica, Sodium Benzoate, Sodium Hyaluronate, Zea Mays (Corn) Starch, Zingiber Officinale (Ginger) Root Extract.

Mycozylal AL

Active ingredient

Tolnaftate 1%

Purpose

Antifungal

Uses

- for the treatment of most athlete's foot (tinea pedis) and ringworm (tinea corporis).
- relieves itching, burning, cracking, scaling, and discomfort which accompany these conditions.
- for the prevention of most Athlete's foot with daily use.
- eliminates fungus on fingers, toes, and around the nails.
- eliminates and helps stop the spread of fungal infections on cuticles around nail edges and under the nail tips where reachable with applicator brush.

Warnings

For external use only

Do not use on

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

When using this product

• do not get into eyes

Stop use and ask a doctor

- for athlete's foot and ringworm if irritation occurs or there is no improvement within 4 weeks
- for prevention of athlete's foot if irritation occurs, discontinue use and consult a doctor

Keep out of reach of children.

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

Directions

- clean the affected area and dry thoroughly
- apply a thin layer of the product over affected area twice daily (morning and night)
 paying special attention to the edges of the nail, cuticles, and skin around the nails 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, ventilated shoes, and change shoes and socks at least once daily
- for athlete's foot and ringworm, use daily for 4 weeks
- if condition persists longer, consult a doctor.

Use under the direction of a medical practitioner

Other information

- Store at 15 30°C (59° 86°F) [see USP Controlled Room Temperature].
- avoid excessive heat
- do not use if package is damaged

How Supplied

Mycozyl AL™ Antifungal Liquid is supplied in a 1 fl oz / 30 mL glass bottle with a screw cap fitted with a brush applicator (NDC 59088-443-03)

Inactive ingredients

Apple Cider Vinegar, Argania Spinosa (Argan) Kernel Oil, Benzyl Alcohol, DMSO (Dimethyl Sulfoxide), Eucalyptus Globulus (Eucalyptus) Leaf Oil, Glycerin, Laureth-4, Lavandula Angustifolia (Lavender) Oil, Melaleuca Alternifolia (Tea Tree) Leaf Oil, PEG-8, DL-alphatocopheryl acetate.

Miconatate[™]

Packaged in the USA by:

PureTek Corporation

Panorama City, CA 91402 For questions or information call toll-free: **877-921-7873**

NDC 59088-778-00

Miconatate™

Mycozyl AP™

Miconazole Nitrate 2% Antifungal Powder (3 oz / 85 g)

Mycozyl AL™

Tolnaftate 1% Antifungal Liquid (1 fl oz / 30 mL)

See enclosed insert(s) inside packaging for product information.

Store at 15 - 30°C (59 - 86°F) [see USP Controlled Room Temperature].

Avoid excessive heat.

Keep this and all medication out of reach of children.

Use under the direction of a medical practitioner

Packaged in the USA by: PureTek Corporation Panorama City, CA 91402 For questions or information call toll-free: 877-921-7873





MICONATATE

miconazole nitrate and tolnaftate kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:59088-778

ı	Packaging	Packaging			
	# Item Co	ode	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:59088	3-778- 1 in 1 Produ	. CARTON; Type 0: Not a Combination act	09/02/2021	

Quantity of Parts Part # Package Quantity Total Product Quantity Part 1 0 BOTTLE, WTH APPLICATOR 1 mL in 30 Part 2 0 BOTTLE 1 g in 85

Part 1 of 2

MYCOZYL AL

tolnaftate liquid

Product Information

Item Code (Source) NDC:59088-443

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV) TOLNAFTATE 0.3 g in 30 mL

Inactive Ingredients		
Ingredient Name	Strength	
TEA TREE OIL (UNII: VIF565UC2G)		
GLYCERIN (UNII: PDC6A3C0OX)		
APPLE CIDER VINEGAR (UNII: 0UE22Q87VC)		
LAURETH-4 (UNII: 6HQ855798J) BENZYL ALCOHOL (UNII: LKG8494WBH)		
DIMETHYL SULFOXIDE (UNII: YOW8V9698H)		
EUCALYPTUS OIL (UNII: 2R040NI662)		
LAVENDER OIL (UNII: ZBP1YXW0H8)		

POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
ARGAN OIL (UNII: 4V59G5UW9X)	

l	P	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1	NDC:59088- 443-03	30 mL in 1 BOTTLE, WTH APPLICATOR; 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	06/09/2021		

Part 2 of 2

MYCOZYL AP

miconazole nitrate powder

Product Information		
Item Code (Source)	NDC:59088-442	
Route of Administration	TOPICAL	

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE - UNII:7NNO0D7S5M)	MICONAZ OLE NITRATE	2 g in 85 g	

Inactive Ingredients		
Ingredient Name	Strength	
LEVOMENOL (UNII: 24WE03BX2T)		
CARTHAMUS TINCTORIUS (SAFFLOWER) OLEOSOMES (UNII: 9S60Q72309)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
GINGER (UNII: C5529G5JPQ)		
CANANGA ODORATA FLOWER (UNII: 76GTF6Z97M)		
XYLITOL (UNII: VCQ006KQ1E)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
NYLON-12 (UNII: 446U8J075B)		
STARCH, CORN (UNII: O8232NY3SJ)		
KUKUI NUT OIL (UNII: TP11QR7B8R)		
HYALURONATE SODIUM (UNII: YSE9PPT4TH)		

I	Packaging					
7	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
:	NDC:59088-442- 07	85 g in 1 BOTTLE; 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	12/08/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	09/02/2021	

Labeler - PureTek Corporation (785961046)

Revised: 10/2024 PureTek Corporation