

GENRX DAILY DEFENSE ANTIFUNGAL- miconazole nitrate powder

PureTek Corporation

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.

Active ingredient

Miconazole Nitrate 2.0%

Purpose

Antifungal

Uses

- for the treatment of most athlete's foot (tinea pedis), jock itch (tinea cruris), ringworm (tinea corporis)
- relieves itching, scaling, cracking, burning, redness, soreness, irritation, discomfort and chafing associated with jock itch

Warnings

For external use only

Do not use on

children under 2 years of age unless directed by a doctor

When using this product

avoid contact with the 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

Inactive ingredients

Aleurites moluccana seed oil, *Aloe barbadensis* (*Aloe vera*) leaf juice, fragrance, GenRx Complex™ [consisting of: bisabolol, calcium pantothenate (vitamin B₅), *Carthamus tinctorius* (safflower) oleosomes, maltodextrin, niacinamide (vitamin B₃), pyridoxine HCl (vitamin B₆), silica, sodium ascorbyl phosphate (vitamin C), sodium starch octenylsuccinate, tocopheryl acetate (vitamin E), *Zingiber officinale* (ginger) root extract], nylon-12, sodium benzoate, sodium hyaluronate, *Zea mays* (corn) starch.

Daily Defense Antifungal Powder with GenRx Complex™ (3 oz bottle label)


Room No.

Name


Drug Facts	
Active ingredient	Purpose
Miconazole Nitrate 2.0%	Antifungal
Uses	
<ul style="list-style-type: none"> ■ for the treatment of most athlete's foot (tinea pedis), jock itch (tinea cruris), ringworm (tinea corporis) ■ relieves itching, scaling, cracking, burning, redness, soreness, irritation, discomfort and chafing associated with jock itch 	
Warnings	
For external use only	
Do not use on children under 2 years of age unless directed by a doctor	
When using this product avoid contact with the eyes	
Stop use and ask a doctor if	
<ul style="list-style-type: none"> ■ for athlete's foot and ringworm - irritation occurs or if there is no improvement within 4 weeks ■ for jock itch - irritation occurs or if 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	
<ul style="list-style-type: none"> ■ 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.	
Inactive ingredients <i>Aleurites moluccana</i> seed oil, <i>Aloe barbadensis</i> (<i>Aloe vera</i>) leaf juice, fragrance, GenRx Complex™ (consisting of: bisabolol, calcium pantothenate (vitamin B ₅), <i>Carthamus tinctorius</i> (safflower) oleosomes, maltodextrin, niacinamide (vitamin B ₃), pyridoxine HCl (vitamin B ₆), silica, sodium ascorbyl phosphate (vitamin C), sodium starch octenylsuccinate, tocopheryl acetate (vitamin E), <i>Zingiber officinale</i> (ginger) root extract), nylon-12, sodium benzoate, sodium hyaluronate, <i>Zea mays</i> (corn) starch.	

PureTek Guarantee: Every PureTek product is manufactured and tested to meet or exceed all USP standards.

NDC 59088-924-07



WOUND CARE
Hypo-Allergenic




Daily Defense
Antifungal Powder

with
GenRx Complex™

Talc-Free Formula
Helps relieve fungal symptoms of
itching, burning and irritation

- Physician Tested
- Non-sensitizing
- Promotes Healing
- Clinically Proven
- Helps Skin Cell Renewal
- Paraben Free

3 oz / 85 g



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LATEX-FREE

Manufactured in the USA by: PureTek Corporation, San Fernando, CA 91340
For more info, visit: www.genrxwoundcare.com • 877-921-7873
LIST NO. 92407AAA Rev: 240271-00

GENRX DAILY DEFENSE ANTIFUNGAL

miconazole nitrate powder

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59088-924
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE - UNII:7NNO0D7S5M)	MICONAZOLE NITRATE	20 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
KUKUI NUT OIL (UNII: TP11QR7B8R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
LEVOMENOL (UNII: 24WE03BX2T)	
CALCIUM PANTOTHENATE (UNII: 568ET80C3D)	
CARTHAMUS TINCTORIUS SEED OLEOSOMES (UNII: 9S60Q72309)	
MALTO DEXTRIN (UNII: 7CVR7L4A2D)	
NIACINAMIDE (UNII: 25X51I8RD4)	
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM ASCORBYL PHOSPHATE (UNII: 836SJG51DR)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
GINGER (UNII: C5529G5JPQ)	
NYLON-12 (UNII: 446U8J075B)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
STARCH, CORN (UNII: O8232NY3SJ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59088-924-07	85 g in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333C	01/08/2013	

Labeler - PureTek Corporation (785961046)

Establishment

Name	Address	ID/FEI	Business Operations
PureTek Corporation		785961046	manufacture(59088-924) , pack(59088-924) , label(59088-924)

Revised: 1/2013

PureTek Corporation