EVERCARE ANTIFUNGAL TREATMENT SUPER ABSORBENT POWDER- miconazole nitrate powder
PANEFFORT, LLC

Evercare Antifungal Treatment Super Absorbent Powder

Drug Facts

Active Ingredient

Miconazole Nitrate 2%

Purpose

Antifungal

Use

for the cure of most athlete's foot (tinea pedis), jock itch (tinea cruris) and ringworm (tea corporis) for effective relief of itching, cracking, burning, scaling and discomfort

Warnings

For external use only.

Do not use while smoking or near heat or flame. **Flammable,**

Do not use

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

When using this product

- avoid contact with eyes. If product gets into eyes, rinse thoroughly with water.

Stop use and ask a doctor

if irritation occurs or there is no improvement within 4 weeks for athlete's foot and ringworm, or 2 weeks for jock itch

Keep out of reach of children.

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

Directions

clean the affected area& dry thoroughly apply a thin layerof 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 attenting to spaces between the

toes;wear wellfitting, ventilated shoes, & change shoes& socks at least once daily for athlete's foot&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

store between 59-86 °F

- lightly shake bottle to loosen settled powder

Inactive ingredients

corn starch, sodium bicarbonate, kaolin, zinc oxide, calamine

Package Labeling:

EVERCARE ANTIFUNGAL TREATMENT SUPER ABSORBENT POWDER

miconazole nitrate powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83099-000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE -	MICONAZOLE NITRATE	20 mg in 1 g

UNII:7NNO0D7S5M)	MICONAZOLE NITRATE	20 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
KAOLIN (UNII: 24H4NWX5CO)	
ZINC OXIDE (UNII: SOI2LOH54Z)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83099-000-00	85 g in 1 POUCH; Type 0: Not a Combination Product	11/28/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	11/28/2022	

Labeler - PANEFFORT, LLC (018250858)

Revised: 10/2023 PANEFFORT, LLC