MICONAZOLE NITRATE- miconazole nitrate powder Westminster Pharmaceuticals, LLC

Miconazole Nitrate

Drug Facts

Active ingredient

Miconazole Nitrate 2.0%

Purpose

Antifungul

Uses

- for the treatment of most athlete's foot (tinea pedis), jock itch (tinea cruris), ringworm (tinea corporis)
- for the treatment of most superficial skin infections caused by yeast (candida albicans)
- relieves most 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 by a doctor
- avoid contact with the eyes
- for athlete's foot and ringworm if irritation occurs, or if there is no improvement within 4 weeks, discontinue use and consult a doctor
- for jock itch if irritation occurs, or if there is no improvement within 2 weeks, dicontinue 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 powder 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 toes; wear well fitting, ventilated shoes, and change shoes and socks at least once daily
- for athletes 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

allantoin, chloroxylenol, fragrance, imidazolidinyl urea, microcrystalline cellulose, tricalcium phosphate, corn starch

PRINCIPAL DISPLAY PANEL - 85 g Bottle Label

NDC 69367-399-85

Miconazole

Antifungul Powder Treatment

Miconazole Nitrate 2%

Botanical Nutrition For Sensitive Skin

- CHG Compatible
- Paraben Free
- Hypoallergenic

NET WT. 3 OZ (85g)

Westminster Pharmaceuticals



	HUMAN OTC DRUG TOPICAL	ltem Code (Source)	NDC:69367-399
Product Type		ltem Code (Source)	NDC:69367-399
Product Type		ltem Code (Source)	NDC:69367-399
		Item Code (Source)	NDC:69367-399
Route of Administration	TOPICAL		
Active Ingredient/Active I	Maiaty		
Ingre	f Strength Strength		
MICONAZOLE NITRATE (UNII: VW4)	H1CYW1K) (MICONAZOLE -	UNII:7NNO0D7S5M) MICONAZ	DLE NITRATE 2 g in 100 g
Inactive Ingredients			
	Ingredient Name		Strength
ALLANTOIN (UNII: 344S277G0Z)			
CHLOROXYLENOL (UNII: 0F32U78V			
IMIDUREA (UNII: M629807ATL)			
MICROCRYSTALLINE CELLULOSE	(UNII: OP1R32D61U)		
TRICALCIUM PHOSPHATE (UNII: K	4C08XP666)		
STARCH, CORN (UNII: 08232NY3SJ))		

Ρ	roduct Chara	icteristics							
Color Shape Flavor		WHITE (White to off-white)		Score					
			Size Imprint Code						
С	ontains								
D	ackaging								
	ackaying								
#	ltem Code	Package Description	Ma	rketing Start Date	Marketing Date	End			
1	NDC:69367-399- 85	85 g in 1 BOTTLE; Type 0: Not a Combination Product	05/16/	2024					
	larketing	Information							
M			м	arketing Start	Marketing	End			
Μ	Marketing Category	Application Number or Monograph Citation		Date	Date				

Labeler - Westminster Pharmaceuticals, LLC (079516651)

Revised: 5/2024

Westminster Pharmaceuticals, LLC