1236-CS ANTIFUNGAL POWDER (TALC FREE)- 1236-ccantifungal powder(talc free) powder
Dynarex Corporation

1236-CS Antifungal Powder (Talc Free)

Active Ingredient

Miconazole Nitrate 2% USP

Purpose

Antifungal

Use(s)

• For the treatment of most athlete's foot (tinea pedis), jock itch (tinea crusis), and ringworm (tinea corporis)

• Relieves itching, scaling, cracking, burning, redness, soreness, irritation, discomfort, and chafing associated with jock itch and itching or burning feet.

Warnings

For External Use Only

Do not use

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

• For diaper rash

Ask a doctor or pharmacist

Before use if you are taking anticoagulants such as warfarin

When using this product

• Avoid contact with the eyes

Stop use and ask a doctor if

• Irritation occurs

• There is no improvement within 2 weeks when used for the treatment of jock itch

• There is no improvement within 4 weeks when used for athlete's foot or ringworm

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222)

right away.

Directions

• Clean the affected area and dry thoroughly. Apply a thin layer over the 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

• Store at Controlled Room Temperature • Do not use if "Remove before Use" seal is missing

Inactive Ingredients

Colloidal Silicon Dioxide, Corn Starch, Dipotassium Glycyrrhizinate, Sodium Benzoate

Questions?

1-888-396-2739 Monday - Friday 9AM - 5PM EST

Label

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1236-ccantifungal powder(talc free) powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-313
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**
STARCH, CORN (UNII: O8232NY3SJ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
GLYCYRRHIZINATE DIPOTASSIUM (UNII: CA2Y0FE3FX)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Product Characteristics			
Color		**Score**	
Shape	FREEFORM	**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:67777-313-04	24 in 1 CASE	04/04/2025	
1	NDC:67777-313-03	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	04/04/2025	

Labeler - Dynarex Corporation (008124539)

Revised: 3/2025 Dynarex Corporation