ZEASORB JOCK ITCH- miconazole nitrate powder
Crown Laboratories

Zeasorb Jock Itch

Active ingredient

Miconazole nitrate 2%

Purpose

Antifungal

Use

- Proven clinically effective in the treament of most jock itch

Warnings

For external use only.

Avoid contact with the eyes.

Do not use

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

If irritation occurs or if there is no improvement within 2 weeks, 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

Wash 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. 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 20 º - 25 ºC (68 º - 77 ºF) [see USP Controlled Room Temperature]. Product settles during shipment. Package contains full net weight.

Inactive ingredients

Aloe Barbadensis Leaf Juice Powder, Bentonite, Fragrance, Potassium Sorbate, Sodium Benzoate, Tapioca Starch, Tricalcium Phosphate, Water, Zea Mays (Corn) Starch

Questions?

call **1-833-279-6522**

Principal Display

New & Improved

Zeasorb®AF

Antifungal Powder

with 2% Miconazole Nitrate

Cures Most Jock Itch

Relieves Itching, Burning, Scaling

Patented Odor Control Technology

Attacks and Absorbs Moisture

Talc-Free, Paraben-Free, Aluminum-Free

Dermatologist Recommended

Net wt. 2.5 oz (71g)

P12322.00

ZEASORB JOCK ITCH

miconazole nitrate powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0316-8001
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**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

STARCH, TAPIOCA (UNII: 24SC3U704I)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
WATER (UNII: 059QF0KO0R)	
STARCH, CORN (UNII: O8232NY3SJ)	
BENTONITE (UNII: A3N5ZCN45C)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0316-8001-02	71 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/19/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	10/19/2023	

Labeler - Crown Laboratories (079035945)

Revised: 11/2023 Crown Laboratories