ZEASORB- miconazole nitrate powder
Crown Laboratories

Zeasorb

Active ingredient

Miconazole nitrate 2%

Purpose

antifungal

Use (Athlete's Foot)

for the cure of most athlete's foot

Use (Jock Itch)

for the cure 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.

Stop use and ask a doctor if (Athlete's Foot)

- irritation occurs or there is no improvement within 4 weeks.

Stop use and ask a doctor if (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 (Athlete's Foot)

- 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
- pay special attention to spaces between the toes; wear well-fitting, ventilated shoes, and change shoes and socks at least once daily
- use daily for 4 weeks

- if condition persists longer, consult a doctor
- this product is not effective on the scalp or nails

Directions (Jock Itch)

- 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
- 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

Aldioxa, Chloroxylenol, Croscarmellose Sodium, Fragrance, Imidazolidinyl Urea, Magnesium Stearate, Microcrystalline Cellulose, Sodium Acrylates Crosspolymer-2

Questions or comments?

call 1-833-279-6522

Distributed by: Crown Laboratories, Inc., Johnson City, TN 37604

ZEASORB is a registered trademark of Crown Laboratories, Inc.

Principal Display Panel

NDC 0316-0231-25

Zeasorb ® AF

ANTIFUNGAL TREATMENT

Miconazole Nitrate 2%

Super Absorbent Powder

New & Improved

cures most **athlete's foot**

Absorbs & Repels Moisture

Relieves:

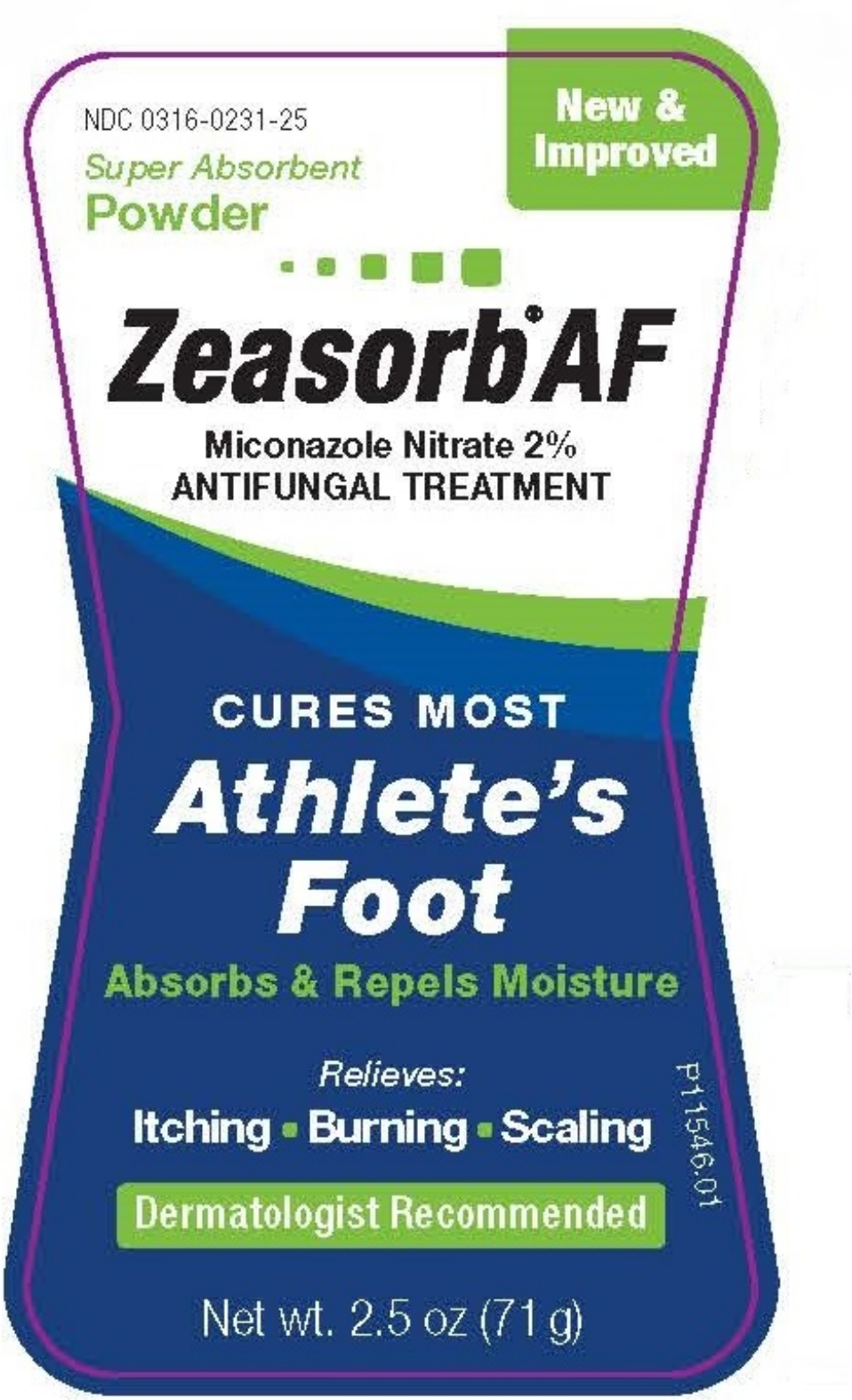
- Itching

- Burning
- Scaling

Dermatologist Recommended

Net wt. 2.5 oz (71 g)

Front Label: P11546.01



Principal Display Panel

NDC 0316-0232-25

Zeasorb ® AF

ANTIFUNGAL TREATMENT

Miconazole Nitrate 2%

Super Absorbent Powder

New & Improved

cures most **jock itch**

Absorbs & Repels Moisture

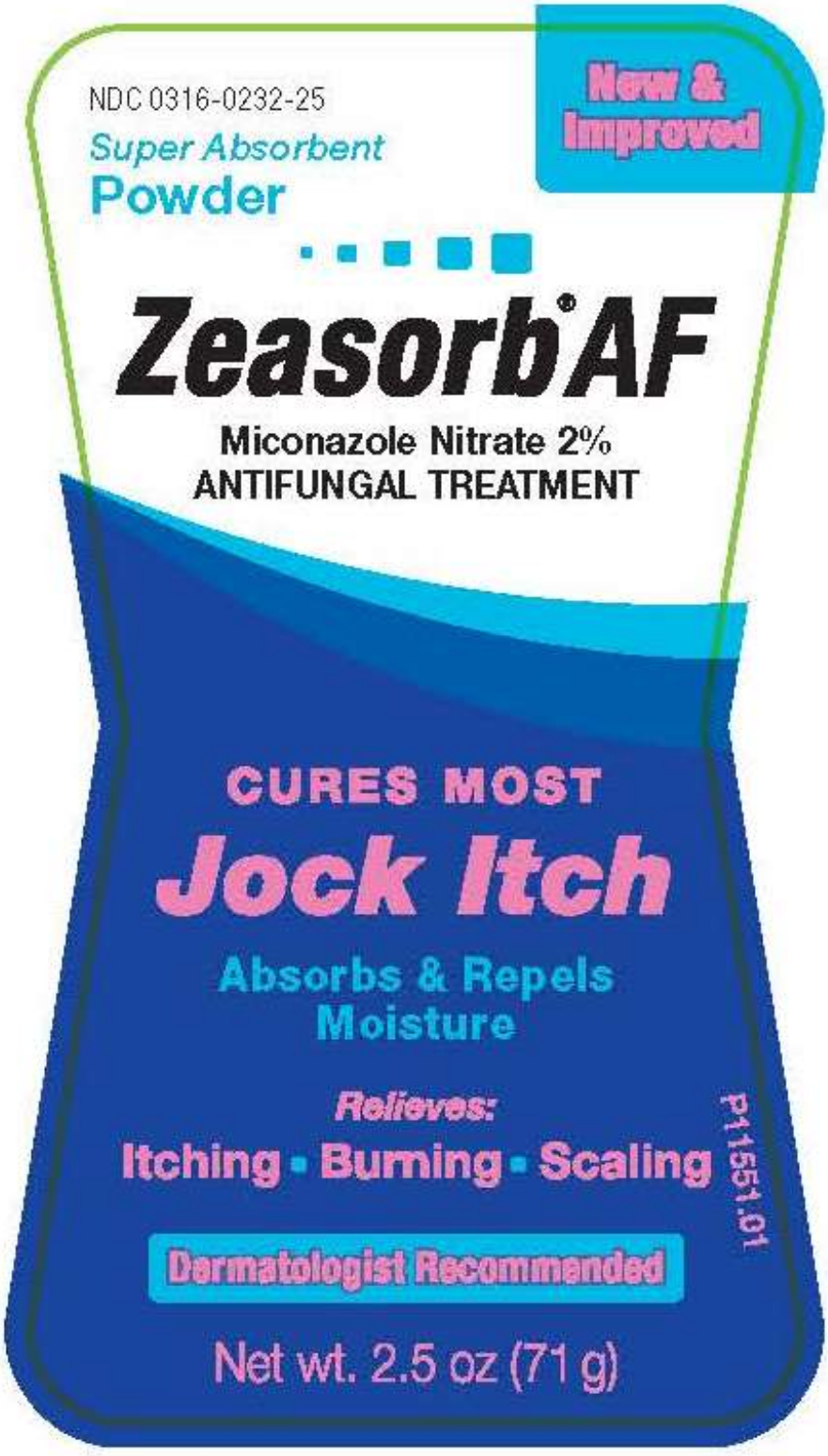
Relieves:

- Itching
- Burning
- Scaling

Dermatologist Recommended

Net wt. 2.5 oz (71 g)

Front Label: P11551.01



ZEASORB

miconazole nitrate powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0316-0232
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
SODIUM ACRYLATES CROSSPOLYMER-2 (UNII: D3HPR4WW6F)	
ALDIOXA (UNII: 8T66I31YNK)	
CHLOROXYLENOL (UNII: 0F32U78V2Q)	
IMIDUREA (UNII: M629807ATL)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	

Product Characteristics

Color	white (white to off-white)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0316-0232-25	71 g in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2018	04/16/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	03/14/2011	04/16/2025

ZEASORB

miconazole nitrate powder

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0316-0231
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
ALDIOXA (UNII: 8T66I31YNK)	
CHLOROXYLENOL (UNII: 0F32U78V2Q)	
IMIDUREA (UNII: M629807ATL)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM ACRYLATES CROSSPOLYMER-2 (UNII: D3HPR4WW6F)	

Product Characteristics

Color	white (white to off-white)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0316-0231-25	71 g in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2018	03/31/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	03/14/2011	03/31/2025

Labeler - Crown Laboratories (079035945)

Revised: 12/2023 Crown Laboratories