

CLOTRIMAZOLE- clotrimazole solution
Akron Pharma Inc

Clotrimazole Topical Solution USP,1%

Drug Facts

Active ingredient

Clotrimazole USP, 1%

Purpose

Antifungal

Uses

Cures most

- Athlete's foot (tinea pedis)
- Jock Itch (tinea cruris)
- Ringworm (tinea corporis)

Effectively relieves

- Itching
- Cracking
- Burning

Discomfort which can accompany these conditions

Warnings

For external use only

Ask a doctor before use

on children under 2 years of age

When using this product

avoid contact with eyes

Stop use and ask a doctor if

- irritation occurs
- 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 a Poison Control Center right away.

Directions

- This product is not effective on the scalp or nails For best results, follow directions and continue treatment for length of time indicated. For athlete's foot and ringworm: use daily for 4 weeks. For jock itch: use daily for 2 weeks.
- 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

For athlete's foot: pay special attention to spaces between the toes; wear well-fitting, ventilated shoes; change shoes and socks at least once daily

Other information

- **Store at 15° to 30°C (59° to 86°F)**

Inactive ingredient

polyethylene glycol 400

Questions?

Please Call **1-877-225-6999**

PRINCIPAL DISPLAY PANEL - 10 mL Bottle Carton

NDC 71399-0500-1

Compare to the active ingredient of Lotrimin[®] AF*

Clotrimazole Topical Solution USP, 1%**Antifungal**

- *Relieves Itching & Burning*
- *Greaseless, Nonstaining*

For External use only.

Not for Ophthalmic use.

Keep this and all medications out of the reach of children .

Akron Pharma

10ml

Drug Facts (continued)**Directions**

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Drug Facts (continued)**Other Information**

- Store between 15°-30°C (59°-86°F)

Inactive Ingredients
polyethylene glycol 400

Manufactured for:
Akron Pharma, Inc., Fairfield, NJ 07004

Compare to the active ingredient of Lotrimin*

10ml

NDC: 71399-0500-1

Clotrimazole (Topical Solution) USP, 1% Anti fungal

- Relieves Itching & Burning
- Greaseless, Non-staining

For External use only,
Not for Ophthalmic use.

Keep this and all medications
out of the reach of children.

Drug Facts

Active Ingredient Clotrimazole USP, 1%.....
Purpose Antifungal

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- Jock Itch (tinea cruris),
- Ringworm (tinea corporis)

Effectively relieves

- itching ■ cracking ■ burning
- Discomfort which can accompany these conditions

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on children under 2 years of age
- When using this product**
avoid contact with eyes
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 - 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 a Poison Control
center right away.

Questions?
Please call 1-877-225-6999

* This product is not manufactured by Shering-Plough Healthcare Products Inc., owner of the registered trademark Lotrimin® AF

Akron Pharma

NDC 71399-0500-3

Compare to the active ingredient of Lotrimin® AF*

Clotrimazole Topical Solution USP, 1%

Antifungal

Relieves Itching & Burning

Greaseless, Nonstaining

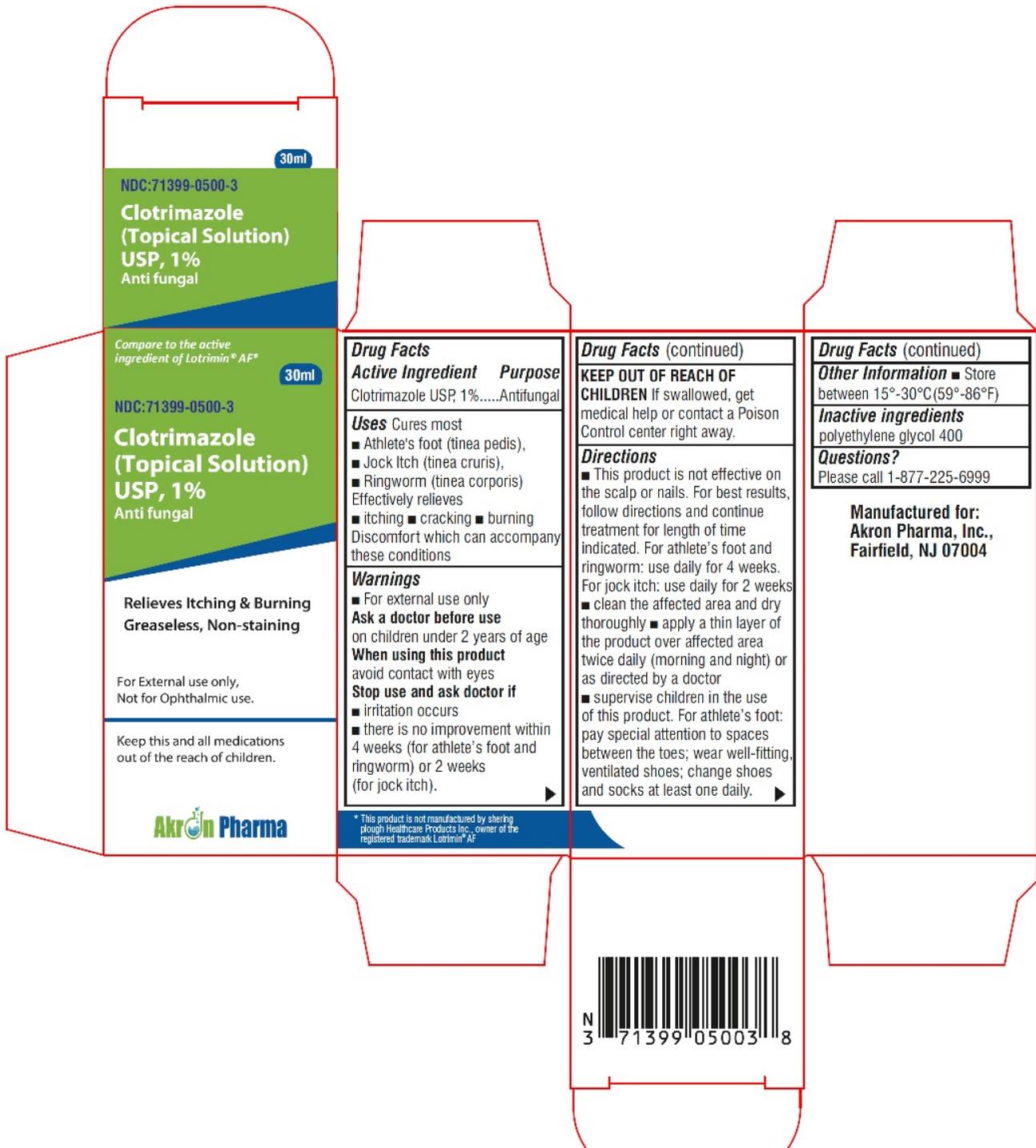
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Akron Pharma

30ml



CLOTRIMAZOLE

clotrimazole solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71399-0500
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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CLOTRIMAZOLE (UNII: G07GZ97H65) (CLOTRIMAZOLE - UNII:G07GZ97H65)	CLOTRIMAZOLE	1 g in 1 mL
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Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-0500-1	10 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/13/2024	
2	NDC:71399-0500-3	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/13/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	09/13/2024	

Labeler - Akron Pharma Inc (067878881)

Revised: 1/2026

Akron Pharma Inc