ANTI-ITCH CREAM- diphenhydramine hydrochloride, zinc acetate cream Kroger

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Kroger Anti-Itch Cream

Drug Facts

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Active ingredients

Diphenhydramine hydrochloride 2%

Zinc acetate 0.1%

Purpose

Topical analgesic

Skin protectant

Uses

-temporarily relieves pain and itching associated with:

- insect bites
- minor burns
- sunburn
- minor skin irritations
- minor cuts
- scrapes
- rashes due to poison ivy, poison oak, and poison sumac

-dries the oozing and weeping of poison ivy, poison oak, and poison sumac

Warnings

For external use only.

Do not use

- on large areas of the body
- with any other product containing diphenhydramine, even one taken by mouth

Ask a doctor before use

- on chicken pox
- on measles

When using this product

avoid contact with eyes

Stop use and ask a doctor if

- condition worsens or does not improve within 7 days
- symptoms persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

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

Directions

- do not use more than directed
- adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: ask a doctor

Other information

- store at 20° - 25°C (68°-77°F)
- protect from excessive heat (40°C/104°F)

Inactive ingredients

cetyl alcohol, diazolidinyl urea, glyceryl stearate/PEG-100 stearate, methylparaben, propylene glycol, propylparaben, purified water

Questions or comments?

Call **1-800-632-6900**

BEND LINE

NO PRINT OR COLOR

NDC 41226-888-02

Extra Strength

Anti-Itch Cream

Diphenhydramine HCl 2%/ Zinc Acetate 0.1%

Topical Analgesic/Skin Protectant

Temporarily relieves pain and itching associated with insect bites; poison ivy, oak, sumac; sunburn; minor cuts & scrapes

NET WT 1 OZ (28 g)

BEND LINE

NO PRINT OR COLOR

NO PRINT OR COLOR

Drug Facts

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Diphenhydramine hydrochloride 2%	Topical analgesic
Zinc acetate 0.1%	Skin protectant

Uses ■ temporarily relieves pain and itching associated with: ■ insect bites ■ minor burns ■ sunburn ■ minor skin irritations ■ minor cuts ■ scrapes ■ rashes due to poison ivy, poison oak, and poison sumac ■ dries the oozing and weeping of poison ivy, poison oak, and poison sumac

Warnings

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Do not use ■ on large areas of the body ■ with any other product containing diphenhydramine, even one taken by mouth

Ask a doctor before use ■ on chicken pox ■ on measles

When using this product avoid contact with eyes

Stop use and ask a doctor if ■ condition worsens or does not improve ▶

Drug Facts (continued)

within 7 days ■ symptoms persist for more than 7 days or clear up and occur again within a few days

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Directions ■ do not use more than directed ■ adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily ■ children under 2 years of age: ask a doctor

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SIGLINE

Questions or comments? Call **1-800-632-6900**

DO NOT USE IF SEAL ON TUBE IS PUNCTURED OR MISSING

PROUDLY DISTRIBUTED BY
THE KROGER CO.
CINCINNATI, OHIO 45202
MADE IN INDIA 809A 1125
QUALITY GUARANTEE
www.kroger.com

NO PRINT OR COLOR

ANTI-ITCH CREAM

diphenhydramine hydrochloride, zinc acetate cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41226-888
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	1 mg in 1 g
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	20 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
WATER (UNII: 059QF0KO0R)	
GLYCERYL STEARATE/PEG-100 STEARATE (UNII: RD25J5V947)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41226-888-02	1 in 1 CARTON	11/15/2025	
1		28 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/15/2025	

Labeler - Kroger (006999528)

Registrant - TIME CAP LABORATORIES, INC. (037052099)

Establishment

Name	Address	ID/FEI	Business Operations
MARKSANS PHARMA LIMITED		677604129	manufacture(41226-888)