

ITCH RELIEF- diphenhydramine hcl, zinc acetate spray
The Kroger Co.

Kroger 295.000/295AA-AB
Itch Relief Spray

Active ingredients

Diphenhydramine HCl 2%

Zinc acetate 0.1%

Purpose

External analgesic

Skin protectant

Uses

- for the temporary relief of pain and itching associated with minor skin irritations
- dries the oozing and weeping of poison: ivy, oak and sumac

Warnings

For external use only

Flammable. Keep away from fire or flame.

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 or measles

When using this product

do not get into eyes

Stop use and ask a doctor if

condition worsens or symptoms last 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.

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

- store at 20° to 25° C (68° to 70°F)

Inactive ingredients

alcohol, glycerin, povidone, purified water, tris (hydroxymethyl) aminomethane

Questions?

1-800-632-6900

Adverse reaction

DISTRIBUTED BY THE KROGER CO. CINCINNATI, OHIO 45202

Disclaimer

Benadryl[®] is a registered trademark of Kenvue, Inc. Kenvue, Inc. is not affiliated with The Kroger Co. or this product.

Principal display panel

TEAR HERE

NDC 59450-295-20

COMPARE TO the active ingredients in BENADRYL[®] EXTRA STRENGTH TOPICAL ANALGESIC SPRAY*

Kroger[®]

Extra Strength

Itch Relief Spray

Diphenhydramine HCl 2%

Topical Analgesic

Zinc Acetone 0.1%

Skin Protectant

Temporary relief of pain and itching due to insect bites, poison ivy or oak, or other minor skin irritations

2 FL OZ (59 mL)

L0021624FA NDC 59450-295-20 **TEAR HERE**
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EXTRA STRENGTH TOPICAL ANALGESIC SPRAY^{*}

Kroger Extra Strength

Itch Relief Spray

Diphenhydramine HCl 2%
Topical Analgesic

Zinc Acetate 0.1%
Skin Protectant

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2 FL OZ (59 mL)

Drug Facts

Active ingredients	Purpose
Diphenhydramine HCl 2%	External analgesic
Zinc acetate 0.1%	Skin protectant

Uses ■ for the temporary relief of pain and itching associated with minor skin irritations ■ dries the oozing and weeping of poison: ivy, oak and sumac

Warnings
For external use only

Flammable. Keep away from fire or flame.

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 or measles

When using this product do not get in eyes

Stop use and ask a doctor if condition worsens or symptoms last 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.

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° to 25° C (68° to 77°F)

Inactive ingredients alcohol, glycerin, povidone, purified water, tris (hydroxymethyl) aminomethane

Questions? 1-800-632-6900

L0021624BA

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*Benadril[®] is a registered trademark of Kenecor, Inc. Kenecor, Inc. is not affiliated with The Kroger Co. or this product.

ITCH RELIEF

diphenhydramine hcl, zinc acetate spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59450-295
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 mL
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	20 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
POVIDONE (UNII: FZ989GH94E)	
TROMETHAMINE (UNII: 023C2WHX2V)	
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59450-295-20	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	02/02/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	02/02/2024	

Labeler - The Kroger Co. (006999528)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091514	manufacture(59450-295)