

ITCH RELIEF- diphenhydramine hcl, zinc acetate spray
Retail Business Services, LLC

Care One 295.000/295AB
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 information

- store at 20° to 25°C (68° to 77°F)

Inactive ingredients

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

Disclaimer

*This product is not manufactured or distributed by Johnson & Johnson Consumer Products Company, distributor of Benadryl[®] Spray.

ADVERSE REACTION

DIST. BY FOODHOLD U.S.A., LLC

LANDOVER, MD 20785

1-877-846-9949 ©2021 Ahold Delhaize Licensing Sari

Quality guaranteed or money back.

principal display panel

TEAR HERE

CAREONE[®]

Itch Relief Spray

TOPICAL ANALGESIC

SKIN PROTECTANT

Pain and Itch Reliever

COMPARE TO THE ACTIVE INGREDIENTS IN BENADRYL[®] SPRAY*

RELIEVES ITCHING DUE TO INSECT BITES, POISON OAK OR IVY, OR OTHER MINOR SKIN IRRITATIONS

OUR PHARMACISTS RECOMMEND

2 FL OZ (59 mL)



3 41520 00600 5

Drug Facts

Active ingredients	Purpose
Diphenhydramine HCl 2%	External anesthetic
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 ■ 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, tri (hydroxymethyl) aminomethane

*This product is not manufactured or distributed by Johnson & Johnson Consumer Products Company, distributor of Benadryl® Spray. DIST. BY FOOEHLD U.S.A., LLC, LANDOVER, MD 20716 1-877-448-8948 ©2021 Abbott Diabetes Livingcare Sdn Bhd Quality guaranteed or your money back. L0019684BA

ITCH RELIEF

diphenhydramine hcl, zinc acetate spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72476-295
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:72476-295-20	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	10/06/2011	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017		10/06/2011	

Labeler - Retail Business Services, LLC (967989935)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment			
Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091514	manufacture(72476-295)

Revised: 2/2026

Retail Business Services, LLC