

**DIPHENHYDRAMINE HCL AND ZINC ACETATE- allergy relief spray aerosol,
spray**

Chain Drug Consortium, LLC

Premier Value Allergy 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 burns
- sunburns
- minor cuts
- scrapes
- insect bites
- minor skin irritations
- 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.

Flammable:

Do not use while smoking or near heat or flame. Do not puncture or incinerate. Contents under pressure. Do not store at temperature above 120°F. Intentional misuse by deliberately concentrating and inhaling contents can be harmful or fatal.

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

Stop use and ask doctor if

- conditions worsens
- symptoms last more than 7 days or clear up and occur again within a few days

When using this product

- avoid contact with eyes
- use only as directed

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- 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 between 20° to 25°C (68° to 77°F)

Inactive ingredients

aloe barbadensis leaf juice, glycerin, purified water, sd alcohol 40-B, tromethamine

Questions?

call 1-866-964-0939

Principal Display Panel

Premier Value

Extra Strength

Allergy Relief Spray

Continuous Spray

Diphenhydramine HCl 2%,

Zinc Acetate 0.1%

- Relieves Itching and Pain from Allergic Dermatitis caused by Insect Bites, Poison Ivy, Sumac and Oak
- Histamine Blocking
- Topical Analgesic
- Skin protectant

Net Wt 2.7 OZ (76 g)

COMPARE TO BENADRYL®



**EXTRA STRENGTH
ALLERGY RELIEF SPRAY
CONTINUOUS SPRAY**

**Diphenhydramine Hydrochloride 2%
Zinc Acetate 0.1%**

- Relieves Itching and Pain from Allergic Dermatitis caused by Insect Bites, Poison Ivy, Sumac and Oak
- Histamine Blocking
- Topical Analgesic
- Skin Protectant



NET WT 2.7 OZ (76g)

Drug Facts

Active ingredients	Purpose
Diphenhydramine HCl 2.0%.....	Topical analgesic
Zinc acetate 0.1%.....	Skin protectant

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

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*This product is not manufactured or distributed by Johnson & Johnson Consumer Products Company, owner of the registered trademark Benadryl®.



DISTRIBUTED BY CHAIN DRUG CONSORTIUM, 3301 NW BOCA RATON BLVD., SUITE 101, BOCA RATON, FL 33431
 MADE IN USA



If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.

DOT 2P M5706

AIRCSLPV-1

DIPHENHYDRAMINE HCL AND ZINC ACETATE

allergy relief spray aerosol, spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-631
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	1.52 g in 76 g
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	0.076 g in 76 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

TROMETHAMINE (UNII: 023C2WHX2V)

ALOE VERA LEAF (UNII: ZY81Z83H0X)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-631-00	76 g in 1 CAN; Type 0: Not a Combination Product	12/10/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	12/10/2010	

Labeler - Chain Drug Consortium, LLC (101668460)

Revised: 2/2024

Chain Drug Consortium, LLC