

**DIPHENHYDRAMINE HCL AND ZINC ACETATE- extra strength itch relief
continuous spray aerosol, spray
Premier Brands of America Inc.**

Premier Brands Extra Strength Itch Relief Continuous Spray

Drug Facts

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:

Keep away from fire or flame. Do not puncture or incinerate. Contents under pressure.
Do not store at temperatures above 120°F

Do not use

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

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?

1-888-287-1915

Principal Display Panel

Premier

Extra Strength

Itch Relief

Continuous Spray

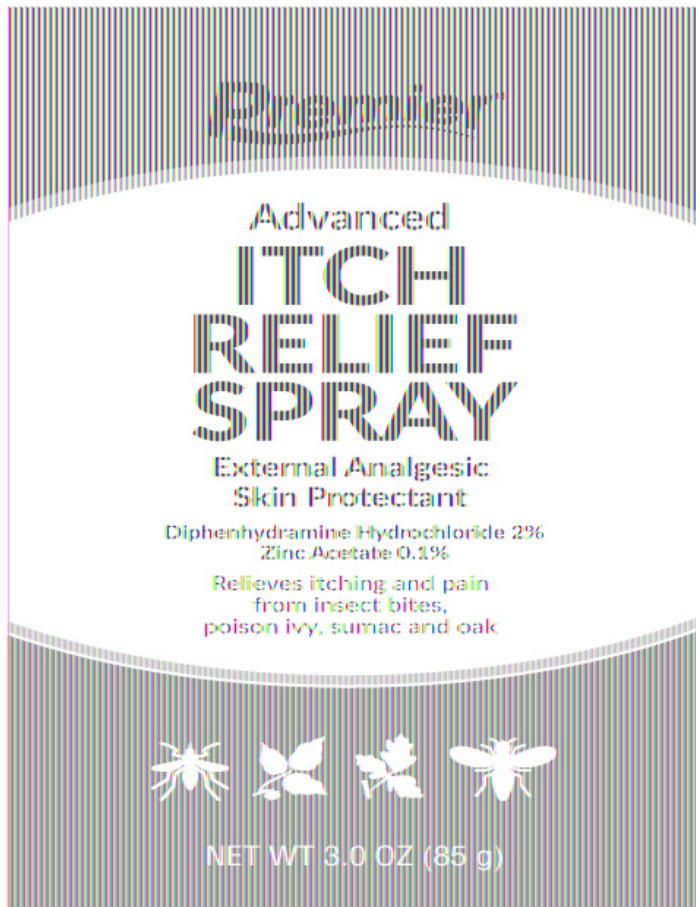
Diphenhydramine HCl 2%,

Zinc Acetate 0.1%

Spray at any angle

- External Analgesic
- Skin Protectant
- Relieve itching and pain from insect bites, rashes due to poison ivy, sumac and oak

3FL OZ (88mL)



Drug Facts

Active ingredients

Diphenhydramine HCl 2.0%
Zinc acetate 0.1%

Uses: for the temporary relief of pain and itching associated with:
 ■ minor burns ■ sunburn ■ minor cuts ■ sore
 ■ minor skin irritations ■ rashes due to poison ivy, poison
 ■ oaks ■ dries the oozing and weeping of poison ivy, poison oak

Warnings

For external use only.

Flammable: Keep away from fire or flame. Do not puncture. Contents under pressure. Do not store at temperatures above 100°F (38°C) to avoid rupture or leakage by deliberately concentrating and inhaling contents.

Do not use: ■ on large areas of the body
 ■ with any other product containing diphenhydramine, even if the other product is for external use only.

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

When using this product: ■ avoid contact with eyes

Stop use and ask doctor if: ■ condition worsens
 ■ symptoms last more than 7 days or clear up and occur again

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

■ do not use more often 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 between 20° to 25°C (68° to 77°F)

Inactive ingredients: aloe barbadensis leaf juice, glycerin, purified water, S/D alcohol 40-B, tromethamine

Questions? Call 1-866-964-0939

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DIPHENHYDRAMINE HCL AND ZINC ACETATE

extra strength itch relief continuous spray aerosol, spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:56104-015
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	1.5 g in 88 mL
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	0.075 g in 88 mL

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:56104-015-03	88 mL in 1 CANISTER; Type 0: Not a Combination Product	01/01/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/01/2013	

Labeler - Premier Brands of America Inc. (117557458)

Revised: 2/2024

Premier Brands of America Inc.