EXTRA STRENGTH ITCH RELIEF- diphenhydramine hcl and zinc acetate spray
Chain Drug Marketing Association Inc

Quality Choice Itch Relief Continuous Spray

Active ingredients

Diphenhydramine HCl 2.0%,

Zinc acetate 0.1%

Purpose

- External analgesic
- Skin protectant

Uses

for the temporary relief of pain and itching associated with:

- minor burns
- sunburn
- 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 120ºF. 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

When using this product

- avoid contact with eyes

Stop use and ask doctor if
- conditions worsens
- 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 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, SD alcohol 40-B, tromethamine

Questions?

call 1-866-964-0939

Principal Display Panel

QC

CONTINUOUS

CHOICE

Extra Strength

Itch Relief

Continous

Spray

Diphenhydramine Hydrochloride 2% /

External Analgesic

Zinc Acetate 0.1% /

Skin Protectant

Relieves itching from insect bites and poison ivy, oak & sumac

Soothes sunburn and minor burns & cuts

NET WT. 2.7 OZ. (76 g)

EXTRA STRENGTH ITCH RELIEF

diphenhydramine hcl and zinc acetate spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-302
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 g

ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	1 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALCOHOL (UNII: 3K9958V90M)	
TROMETHAMINE (UNII: 023C2WHX2V)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-302-27	76 g in 1 CAN; Type 0: Not a Combination Product	06/06/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	06/06/2024	

Labeler - Chain Drug Marketing Association Inc (011920774)

Revised: 6/2024 Chain Drug Marketing Association Inc