ITCH RELIEF- diphenhydramine hcl, zinc acetate spray
Cardinal Health, 110 dba Leader

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Itch Relief Spray
295.000/295AA-AB

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, sumac

Warnings

For external use only

Flammable

Keep away from fire of 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 20º - 25ºC (68º - 77ºF)

Inactive ingredients

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

Questions & comments?

1-800-593-0593

Adverse reactions

Dist. by CAH, Dublin, OH 43017

*This product is not manufactured or distributed by Johnson & Johnson Consumer Products Company of the registered trademark Benadryl Extra Strength Spray.

principal display panel

TEAR HERE

LEADER

NDC 70000-0023-1

Extra Strength

Itch Relief Spray

Diphenhydramine HCl, 2% Zinck Acetate, 0.1%

External Analgesic

Skin Protectant

COMPARE TO BENADRYL EXTRA STRENGTH SPRAY

active ingredients

100% Money Back Guarantee

Relieves Itching

Due to Insect Bits,

Poison Oak or Ivy, or Other Minor Skin Irritations

2 FL OZ (59 mL)

ITCH RELIEF

diphenhydramine hcl, zinc acetate spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70000-0023
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: 059QF0KO0R)	
TROMETHAMINE (UNII: 023C2WHX2V)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70000-0023-1	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/02/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	08/02/2011	

Labeler - Cardinal Health, 110 dba Leader (063997360)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(70000-0023)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(70000-0023)

Revised: 11/2022 Cardinal Health, 110 dba Leader