ITCH RELIEF- diphenhydramine hcl, zinc acetate spray
OLD EAST MAIN CO

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.

Anti-Itch 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

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 worses 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 to 25º C (68º to 77º F)

Inactive ingredients

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

*Not manufactured or distributed by Johnson & Johnson Consumer Products Company, distributor of Benadryl Spray

DISTRIBUTED BY ONE EAST MAIN CO.

100 MISSION RIDE

GOODLETTSVILLE, TN 37072

principal display panel

TEAR HERE

DG health

Compare to active ingredents in Benadryl Spray

Itch Relief Spray

External Analgesic

Skin Protectant

Relieves itching due to insect bites, poison oak or ivy, or other monor skin irritations

100% Satisfaction Guaranteed! (888)309-9030

2 FL OZ (59 mL)

ITCH RELIEF

diphenhydramine hcl, zinc acetate spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-295
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: 059QF0KO0R)	
TROMETHAMINE (UNII: 023C2WHX2V)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-295-20	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/14/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	07/14/2010	

Labeler - OLD EAST MAIN CO (068331990)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(55910-295)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(55910-295)

Revised: 10/2023 OLD EAST MAIN CO