ITCH RELIEF- diphenhydramine hcl, zinc acetate spray
Meijer Distribution, Inc

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

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 or 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 in 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 20° to 25° C (68° to 77° F)

Inactive ingredients

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

*This product is not manufactured or distributed by Johnson & Johnson Consumer Products Company, distributor of Benadryl Spray.

Distributed by Meijer Distribution, Inc.

Grand Rapids, MI 49544

www.meijer.com

295.000/295AB

Principal display panel

Meijer

*Compare to the active ingredients in Benadryl Spray

Itch Relief Spray

topical analgesic

skin protectant

Pain & Itch Reliever

2 FL OZ (59 mL)

ITCH RELIEF

diphenhydramine hcl, zinc acetate spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41250-295
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	18 mg in 1 mL
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	882 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:41250-295-20	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	11/01/2018	

Labeler - Meijer Distribution, Inc (006959555)

Registrant - Vi-Jon (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		790752542	manufacture(41250-295)

Revised: 5/2020 Meijer Distribution, Inc