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.

Meijer 295.000-295AA-AB Itch Relief Spray

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
- on 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 20º to 25º C (68º to 77º F)

Inactive ingredients

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

Disclaimer

*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

principal display panel

TEAR HERE

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
	NDC:41250	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a		

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, LLC (790752542)

Establishment

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

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(41250-295)

Revised: 4/2022 Meijer Distribution, Inc