

ANTI-ITCH WAL-DRYL- diphenhydramine hcl, zinc acetate spray
Walgreen's 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.

Walgreens 295.000/295AA/AB Anti-Itch 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: iv, 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 20° - 25°C (68° - 77°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: Walgreen Co.,
200 Wilmot Rd., Deerfield, IL 60015-4616

100% Satisfaction Guaranteed
www.walgreens.com

principal display panel

TEAR HERE

Compare to Benadryl Spray active ingredient

Walgreens Anti-itch Spray

WAL-DRYL

Diphenhydramine Hydrochloride 2%

Zinc Acetate 0.1%

- Topical Analgesic
- Skin Protectant
- Antihistamine

ITCH RELIEF

2 FL OZ (59 mL)



ANTI-ITCH WAL-DRYL

diphenhydramine hcl, zinc acetate spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-0295
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	2 kg in 100 L
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	0.1 kg in 100 L

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
POVIDONE (UNII: FZ989GH94E)	
WATER (UNII: 059QF0K00R)	
TROMETHAMINE (UNII: 023C2WHX2V)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:0363	0.059 L in 1 BOTTLE, SPRAY, Type 0, Not a		

1	NDC:0363-0295-16	0.059 L in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/08/2011	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC monograph not final	part348		07/08/2011	

Labeler - Walgreen's Co (008965063)

Registrant - Vi-Jon, LLC (790752542)

Establishment			
Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(0363-0295)

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Revised: 12/2022

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