

RITE AID ITCH RELIEF- diphenhydramine hcl 2%, zinc acetate 0.1% spray
Rite Aid

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.

Rite Aid Itch Relief Spray, Diphenhydramine HCl 2%, Zinc Acetate 0.1%

Diphenhydramine HCl 2%, Zinc Acetate 0.1%

Topical Analgesic, Skin Protectant

For the temporary relief of pain and itching associated with insect bites, minor burns, sunburn, minor cuts, scrapes, minor skin irritations, and rashes due to poison ivy, oak, and sumac. Dries the oozing and weeping of poison ivy, oak, and sumac.

For external use only. Flammable--Keep away from fire or flame. **Do not use** on chicken pox, on large areas of the body, with any other products containing diphenhydramine, even one taken by mouth. **When using this product** avoid contact with eyes. In case of contact with eyes, flush thoroughly with water. Contents under pressure. Do not puncture or incinerate. Do not store at temperature above 120F. . **Stop use and ask a doctor if** condition worsens, if symptoms persist for more than 7 days or clear up and occur again within a few days.

If swallowed, get medical help or contact a Poison Control Center right away.

Shake well before use. Adults and children 12 years of age and older, apply to the affected area not more than 3 to 4 times daily. Children under 12 years of age: ask a doctor. To apply to face, spray into palm of hand and gently apply.

Alcohol, Glycerin, PVP, Purified Water, Tromethamine.



ITCH RELIEF SPRAY

DIPHENHYDRAMINE HCl 2%
ZINC ACETATE 0.1%

TOPICAL ANALGESIC & SKIN PROTECTANT

relieves itching & pain due to:
insect bites, poison ivy,
poison oak, poison sumac



2 FL OZ (59 mL)

Drug Facts

**DO NOT USE IF SAFETY SEAL
LABEL IS TORN OR MISSING**

Active ingredients

Diphenhydramine HCl 2%.....	Topical Analgesic
Zinc Acetate 0.1%.....	Skin Protectant

Purpose

Uses For the temporary relief of pain and itching
associated with • rashes due to poison ivy oak and sumac

associated with - rashes due to poison ivy, oak and sumac

- insect bites
- minor skin irritations
- minor cuts
- dries the oozing and weeping of poison ivy, oak and sumac

Warnings

Flammable - Keep away from fire or flame

For external use only

Do not use • on chicken pox • on large areas of the body • with any other products containing diphenhydramine, even one taken by mouth

When using this product • avoid spraying in eyes

Stop use and ask a doctor if

TEAR HERE

- condition worsens
- rash or irritation develops and lasts for more than 7 days or clears up and occurs again within a few days

Keep out of the reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away

Directions • adults and children 2 years or 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 15°-30° C (59° - 86°F)

Inactive ingredients

alcohol, glycerin, PVP, purified water, tromethamine

**This product is not manufactured or distributed by Johnson & Johnson Consumer Inc. owner of the registered*



Consumer Inc, owner of the registered trademark Benadryl®

DISTRIBUTED BY: RITE AID,
30 HUNTER LANE,
CAMP HILL, PA 17011
www.riteaid.com

SATISFACTION GUARANTEE:
If you're not satisfied, we'll
happily refund your money.



RITE AID ITCH RELIEF

diphenhydramine hcl 2%, zinc acetate 0.1% spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-0647
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
TROMETHAMINE (UNII: 023C2WHX2V)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
POVIDONE (UNII: FZ989GH94E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-0647-3	85 g in 1 CAN; Type 0: Not a Combination Product	03/31/2021	

2	NDC:11822-0647-2	59 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/31/2021	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC monograph not final	part348		03/31/2021	

Labeler - Rite Aid (014578892)

Registrant - Derma Care Research Labs, LLC (116817470)

Revised: 7/2022

Rite Aid