

WARRIOR FIRST AID ANTI ITCH- diphenhydramine hcl 2%, zinc acetate

0.1% spray

Derma Care Research Labs, LLC

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.

Warrior First Aid Anti Itch 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.

Do not use more than directed. Adults and children 2 years of age and older, apply to the affected area not more than 3 to 4 times daily. Children under 2 years of age: ask a doctor.

Glycerin, PVP, SD Alcohol, 40, Tromethamine, Water.





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MANUFACTURED FOR UNITED SPIRIT OF AMERICA, INC. PEACHTREE CITY, GA 30269	DAPA SP0200-11-H0034 CARDINAL HEALTH SUP 117301 OWENS & MINOR VENDOR 6516
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Drug Facts DO NOT USE IF SAFETY SEAL LABEL IS TORN OR MISSING

Active ingredient Diphenhydramine HCl 2% ... Zinc Acetate — 0.1%	Purpose Topical analgesic Skin Protectant
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Uses temporarily relieves itching and pain associated with minor irritations, burns, scrapes, cuts, insect

TEAR HERE

Drug Facts (continued)
 bites and rashes due to poison ivy, poison oak & poison sumac. Dries the oozing and weeping of poison ivy, poison oak & poison sumac.

Warnings
 For external use only.
Flammable. Keep away from fire or flame.
Do not use • with any other product containing diphenhydramine, even one taken by mouth • on large areas of the body • on chicken pox • on measles
When using this product • avoid contact with the eyes
Stop use and ask doctor if • condition worsens or does not improve within 7 days • symptoms persist for more than 7 days or clear up and occur again in 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 15° - 30°C (59° - 86°F)
Inactive ingredients Glycerin, PVP, SD Alcohol 40, Tromethamine, Water.

WARRIOR FIRST AID ANTI ITCH

diphenhydramine hcl 2%, zinc acetate 0.1% spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72839-647
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40)	DIPHENHYDRAMINE	2 g

(DIPHENHYDRAMINE - UNII:8GTS82S83M)		HYDROCHLORIDE	in 100 mL	
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)		ZINC ACETATE	0.1 g in 100 mL	
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:72839-647-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/25/2022	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final		part348	07/25/2022	

Labeler - Derma Care Research Labs, LLC (116817470)

Registrant - Derma Care Research Labs, LLC (116817470)

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Derma Care Research Labs, LLC