## FAMILY CARE ANTI-ITCH- diphenhydramine hydrochloride zinc acetate cream United Exchange Corp.

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.

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Active Ingredients

Purpose

Diphenhydramine hydrochloride 1%...... Topical analgesic

Zinc acetate 0.1%...... Skin protectant

Uses

Temporarily relieves pain and itching associated with:

- insect bites
- minor burns
- sunburn
- minor skin irritations
- minor cuts
- scrapes
- rashes due to poison ivy
- poison oak
- poison sumac
- drise the oozing and weeping of poison ivy, poison oak, and poison sumac

#### Warnings

For external use only

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 avoid contact with eyes

Stop use and ask a doctor if

- condition worsens or does not improve within 7 days
- symptoms persist for 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 immediately

#### 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 between 20° to 25°C (68° to 77°F)
- Lot No. & Exp. Date: see box or see crimp of tube

Inactive ingredients

 $cetanol,\,methylparaben,\,polyoxyl\,40\,\,stearate,\,propylene\,\,glycol,\,propylparaben,\,purified\,water,\,stearyl\,alcohol$ 

Distributed by:

United Exchange Corp.

17211 Valley View Ave.

Cerritos CA 90703 USA

Made in Korea



**Temporary Relief** of Pain and Itching Associated with:

- Insect Bites
- Poison Ivy
- Poison Oak
- Poison Sumac

**ITCH RELIEF** 



Compare to the active ingredients of Original Strength BENADRYL®

## Relieves Pain & Itching

Relieves Pain & Itching

- **Temporary Relief** of Pain and Itching **Associated with:**
- Insect Bites
- Poison Ivy
- Poison Oak
- **Poison Sumac**

REGULAR STRENGTH

# ANTI-ITCH Cream

1% DIPHENHYDRAMINE HYDROCHLORIDE USP

NET WT 1 0Z (28 g)





United Exchange Corp. 17211 Valley View Ave. Cerritos, CA 90703 USA www.ueccorp.com Toll Free: 1 800 814 8028

Made in Korea



Inactive ingredients cetanol, methylparaben, polyoxyl 40 stearate, propylene glycol, propylparaben, purified water, stearyl alcohol

Other information a store between 20° to 25°C (68° to 77°F) a Lot No. & Exp. Date: see box or see crimp of tube

DIYECTIONS - do not use more than directed a adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily a children

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

Stop use and ask a doctor if a condition worsens or does not improve within 7 days a symptoms persist for more than 7 days or clear up and occur again within a Mhen using this product avoid contact with eyes

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LOT & EXP

and weeping of poison ivy, poison oak, and poison sumac

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Diphenhydramine hydrochloride 1% Zinc acetate 0.1%.....

Active ingredients

Topical analgesic ....Skin protectant Purpose

Drug Facts



diphenhydramine hydrochloride zinc acetate cream

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65923-495	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
<b>DIPHENHYDRAMINE HYDRO CHLO RIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	1 g in 100 g		
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	.1 g in 100 g		

Inactive Ingredients			
Ingredient Name	Strength		
METHYLPARABEN (UNII: A2I8C7HI9T)			
POLYOXYL 40 STEARATE (UNII: 13A4J4NH9I)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
PROPYLPARABEN (UNII: Z8IX2SC1OH)			
WATER (UNII: 059QF0KO0R)			
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:65923-495-28	1 in 1 CARTON				
1		28 g in 1 TUBE				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part347	11/24/2014		

### Labeler - United Exchange Corp. (840130579)

Revised: 11/2014 United Exchange Corp.