

WALGREENS ANTI ITCH - diphenhydramine hydrochloride, zinc acetate cream
WALGREEN COMPANY

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.

Drug Facts

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, and poison sumac

Dries 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

Directions

- do not use more than directed
- adults and children 2 years of age and older: apply to affected area no more than 3 to 4 times daily
- children under 2 years of age: ask a doctor

Other information

- Store between 20°C and 25°C (68°F and 77°F)

Inactive ingredients

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

DISTRIBUTED BY: WALGREEN CO.

200 WILMOT RD., DEERFIELD, IL 60015

walgreens.com

Made in Korea

Compare to Benadryl®
Original Strength active ingredients*

Walgreens

Anti-Itch Cream

1% Diphenhydramine Hydrochloride USP Original Strength

Itch Stopping Cream

Regular Strength Itch Relief

Walgreens

Anti-Itch Cream

1% Diphenhydramine Hydrochloride USP Original Strength

NET WT 1 OZ (28g)

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Walgreens
**100% SATISFACTION
GUARANTEED**
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ITEM 150620



*This product is not manufactured or distributed by McNeil-PPC, Inc, owner of the registered trademark Benadryl®.

Drug Facts
Active ingredients Diphenhydramine hydrochloride 1% Zinc acetate 0.1% Topical analgesic 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, and poison sumac. Dries the oozing and weeping of poison ivy, poison oak, and poison sumac.
Warnings Do not use ■ on large areas of the body ■ with any other product containing diphenhydramine, even one taken by mouth. For external use only.
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
Directions ■ do not use more than directed ■ adults and children 2 years of age and older: apply to affected area no more than 3 to 4 times daily ■ children under 2 years of age: ask a doctor
Other information ■ Store between 20°C to 25°C (68°F to 77°F)
Inactive ingredients cetanol, methylparaben, polyoxy/40 stearate, propylene glycol, propylparaben, purified water, stearyl alcohol

LOT & EXP.

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WALGREENS ANTI ITCH

diphenhydramine hydrochloride, zinc acetate cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-1411
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	1 g in 100 g
ZINC ACETATE (UNII: FM5526K07A) (ZINC - UNII:J41CSQ7QDS)	ZINC ACETATE	0.1 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: 059QF0K00R)	
STEARYL ALCOHOL (UNII: 2KR89I4HIY)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-1411-01	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	part336	03/01/2012	

Labeler - WALGREEN COMPANY (008965063)

Revised: 3/2012

WALGREEN COMPANY