# QUALITY CHOICE EXTRA STRENGTH ANTI ITCH - diphenhydramine hydrochloride, zinc acetate cream

#### CHAIN DRUG MARKETING ASSOCIATION INC

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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# **Drug Facts**

Active Ingredients Purpose

Diphenhydramine hydrochloride USP, 2% Antihistamine

Zinc acetate, 0.1% Skin protectant

#### Uses

- for the temporary relief of itching and pain associated with minor skin irritations and rashes due to insect bites, 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 chicken pox, measles, sunburn, oozing skin, broken blisters or on extensive areas of the skin more than directed
- any other drug containing diphenhydramine while using this product

When using this product avoid contact with the eyes

Stop use and ask a doctor if condition worsens, or if symptoms persists 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 right away

#### Directions

- adults and children 12 years of age and older, apply to affected area not more than 3 to 4 times daily, or as directed by a doctor
- children under 12 years of age, consult a doctor

#### Other information

- store at room temperature
- Lot No. and Exp. Date: see box or see crimp of tube

# Inactive ingredients

glycerin, sodium chloride, petrolatum, cetyl alcohol, dimethicone, isopropyl palmitate, distearyldimodium chloride (1-Octadecanaminium, N, N-Dimmethyl-N-Octadecyl-Chloride), methylparaben, propylparaben, purified water, hydroxyethyl cellulose

## DISTRIBUTED BY QUALITY CHOICE

NOVI, MI 48376-0995

MADE IN KOREA

QC 94731



# QUALITY CHOICE EXTRA STRENGTH ANTI ITCH diphenhydramine hydrochloride, zinc acetate cream **Product Information** Product Type HUMAN OTC DRUG Item Code (Source) NDC:63868-947

Route of Administration	TOPICAL
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Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
<b>DIPHENHYDRAMINE HYDRO CHLO RIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	2 g in 100 g			
ZINC ACETATE (UNII: FM5526K07A) (ZINC - UNII:J41CSQ7QDS)	ZINC ACETATE	0.1 g in 100 g			

Inactive Ingredients			
Ingredient Name	Strength		
GLYCERIN (UNII: PDC6A3C0OX)			
SO DIUM CHLO RIDE (UNII: 451W47IQ8 X)			
PETROLATUM (UNII: 4T6H12BN9U)			
CETYL ALCOHOL (UNII: 936JST6JCN)			
DIMETHICO NE (UNII: 92RU3N3Y1O)			
ISOPROPYL PALMITATE (UNII: 8 CRQ2TH63M)			
METHYLPARABEN (UNII: A2I8C7HI9T)			
PROPYLPARABEN (UNII: Z8IX2SC1OH)			
WATER (UNII: 059QF0KO0R)			

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:63868-947-30	1 in 1 CARTON			
1	30 g in 1 TUBE			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph final	part336	02/13/2012			

# **Labeler -** CHAIN DRUG MARKETING ASSOCIATION INC (011920774)

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