ANTI ITCH TOPICAL ANALGESIC- diphenhydramine hydrochloride, zinc acetate cream
Chain Drug Marketing Associations 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.

Quality Choice Extra Strength Itch Stopping Cream 1 oz. 94731 TG/CMI 2019

Active Ingredients Purpose

Diphenhydramine HCI 2%.....Topical analgesic

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

Uses

- temporarily relieves pain and itching associated with:
- insect bites
- minor burns
- minor skin irritations
- sunburn
- minor cuts
- scrapes
- rashes due to poison ivy, oak, and sumac
- dries the oozing and weeping of poison:
- ivy oak
- sumac

Warnings

For external use only

Do not use

- with any other product containing diphenhydramine, even one taken by mouth
- on large areas of the body

Ask a doctor before use

- on chicken pox
- on measles

When using this product avoid contact with the 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

If pregnant or breast-feeding, ask a health professional before use.

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 12 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 12 years of age: consult a doctor

Other information

- store at 20° to 25°C (68° to 77°F)

inactive ingredients

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

DISTRIBUTED BY:

C.D.M.A. INC.

43157 W. NINE MILE

NOVA, MI 48376-0995

Made in Korea

ANTI ITCH TOPICAL ANALGESIC

diphenhydramine hydrochloride, zinc acetate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-218

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**
CETYL ALCOHOL (UNII: 936JST6JCN)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYOXYL 40 STEARATE (UNII: 13A4J4NH9I)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63868-218-01	1 in 1 BOX	10/12/2016	
1		28 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	02/20/2014	

Labeler - Chain Drug Marketing Associations Inc (011920774)

Revised: 12/2019 Chain Drug Marketing Associations Inc