

**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 HCl 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



NDC 63868-218-01

\*Compare to the active ingredients in BENADRYL®

# Extra Strength Anti-Itch Cream

Topical Analgesic - Itching & Pain Relief

Diphenhydramine Hydrochloride 2% | Antihistamine  
Zinc Acetate 0.1% | Skin Protectant

Temporary Relief of Itching & Pain from Minor Skin Irritations & Rashes due to Insect Bites, Poison Ivy, Oak & Sumac

# Extra Strength Anti-Itch Cream

Topical Analgesic - Itching & Pain Relief



1 oz. (28 g) Net Weight



Extra Strength Anti-Itch Cream



Distributed by C.D.M.A., Inc.®  
43157 W. Nine Mile  
Novi, MI 48376-0995  
www.qualitychoice.com  
Questions: 248-449-9300

**TAMPER-EVIDENT: Do not use if "FOIL SEAL" is broken or missing.**

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

MADE IN KOREA

LOT & EXP.

**Drug Facts**

**Active Ingredients**  
Diphenhydramine hydrochloride USP, 2%.....  
Zinc acetate, 0.1%.....  
.....Topical analgesic  
.....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 large areas of the body ■ with any other product containing diphenhydramine, even one taken by the mouth  
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  
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, or as directed by a doctor  
■ children under 12 years of age, consult a doctor

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

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

**Purpose**  
.....Topical analgesic  
.....Skin protectant

## 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)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
POLYOXYL 40 STEARATE (UNII: 13A4J4NH9I)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYL PARABEN (UNII: Z8IX2SC1OH)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
WATER (UNII: 059QF0K00R)	

### 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