

CAREALL ANTI-ITCH- diphenhydramine hydrochloride and zinc acetate cream cream

New World Imports, Inc

Active Ingredients

Diphenhydramine Hydrochloride 2%

Zinc Acetate 0.1%

Purpose

Diphenhydramine Hydrochloride 2%....Topical Analgesic

Zinc Acetate 0.1%.....Skin Protectant

Keep out of Reach of Children

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

Uses

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.
- dries the oozing and weeping of poison ivy, poison oak and poison sumac.

Warnings

For external use only.

Due 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 persists for more than 7 days or clear up and occur again within a few days

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

Inactive Ingredients

carbomer homopolymer type c, cetyl alcohol, glycerin, glyceryl monostearate, light mineral oil, methylparaben, polysorbate 60, propylparaben, purified water, stearic acid

CAREALL[®]

Extra Strength Itch Relief

COMPARE TO ACTIVE INGREDIENTS OF EXTRA STRENGTH BENADRYL[®] CREAM*

Temporary pain and itch relief for:

- Insect Bites • Minor Burns • Sunburn
- Rashes due to Poison Ivy, Oak & Sumac • & More

Anti-Itch Cream *Fragrance-free
Phthalate-free
Dye-free*

EXTRA STRENGTH

CAREALL[®]

Anti-Itch Cream

EXTRA STRENGTH

DIPHENHYDRAMINE HCL 2% • TOPICAL ANALGESIC
ZINC ACETATE 0.1% • SKIN PROTECANT

**HISTAMINE
BLOCKER**

**NET WT.
1.25 OZ (35g)**

Anti-Itch Cream



EXTRA STRENGTH



This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., Owner of the Registered Trademark Benadryl®
 Distributed by: CareAll Products, 160 Athens Way, Nashville, TN 37228 www.careallproducts.com Made in India NDC # 51824-088-01

Drug Facts		Drug Facts (continued)	
Active Ingredients Diphenhydramine Hydrochloride 2% 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 ■ 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		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 15°- 30°C (59°- 86°F). ■ Lot No. & Exp Date: See box or crimp of tube		Inactive Ingredients Carbomer Homopolymer Type C, Cetyl Alcohol, Glycerin, Glyceryl Monostearate, Light Mineral Oil, Methylparaben, Polysorbate 60, Propylparaben, Purified Water, Stearic Acid	
Purpose 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.		Drug Facts (continued)	



CAREALL ANTI-ITCH

diphenhydramine hydrochloride and zinc acetate cream cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51824-088
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	20 mg in 1 g
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	1 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	

CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51824-088-01	35 g in 1 TUBE; Type 0: Not a Combination Product	07/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	07/01/2022	

Labeler - New World Imports, Inc (075372276)