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:

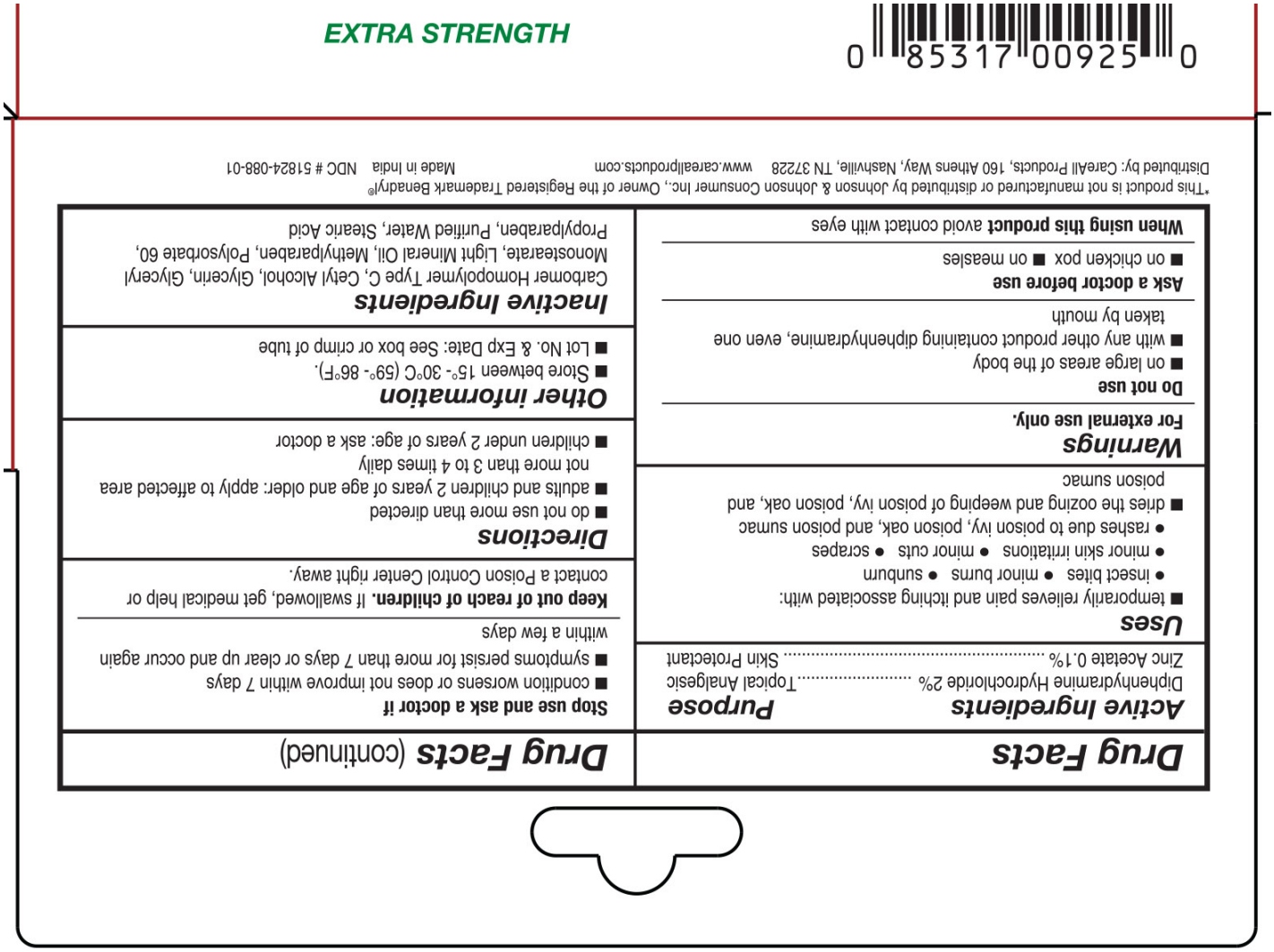
- condition worsens or does not improve within 7 days
- symptoms persists for more than 7 days or clear up and occur again within a few days

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 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)

Revised: 6/2024 New World Imports, Inc