

ANTI ITCH- diphenhydramine hydrochloride, zinc acetate cream
Meijer Distribution Inc

Meijer Distribution, Inc. Anti-Itch Cream Drug Facts

Active ingredients

Diphenhydramine hydrochloride 2%

Zinc acetate 0.1%

Purpose

Topical analgesic

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

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 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 2 years of age and older: apply to affected area no more than 3 to 4 times daily
- children under 2 years of age: ask a doctor

Other information

- store at 20°-25°C (68°-77°F)

Inactive ingredients

cetyl alcohol, diazolidinyl urea, methylparaben, PEG-2 stearate, PEG-20 stearate, propylene glycol, propylparaben, purified water

Questions or comments?

1-800-719-9260

Principal Display Panel

meijer®

EXTRA STRENGTH

Anti-Itch Cream

TOPICAL ANALGESIC | SKIN PROTECTANT

Itch Stopping Cream

Insect Bites | Poison Ivy, Oak, Sumac

Mosquito Bites | Sunburn | Minor Cuts & Scrapes

Fast Relief for Most Outdoor Itches

meijer®

EXTRA STRENGTH

Anti-Itch Cream

COMPARE TO EXTRA STRENGTH BENADRYL® ITCH STOPPING CREAM ACTIVE

INGREDIENTS

TOPICAL ANALGESIC | SKIN PROTECTANT

Itch And Rash Relief

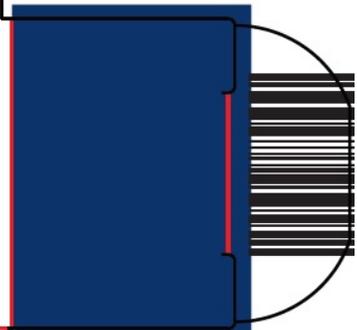
NET WT 1 OZ (28 g)





<p>Drug Facts</p> <p>Active ingredients Diphenhydramine hydrochloride 2%.....Topical analgesic Zinc acetate 0.1%.....Skin protectant</p> <p>Purpose Topical analgesic Skin protectant</p> <p>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</p> <p>Warnings For external use only</p> <p>Do not use ■ on large areas of the body ■ with any other product containing diphenhydramine, even one taken by mouth</p> <p>Ask a doctor before use ■ on chicken pox ■ on measles</p> <p>When using this product ■ avoid contact with the eyes ▶</p>	<p>Drug Facts (continued)</p> <p>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</p> <p>Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.</p> <p>Directions ■ do not use more than directed ■ adults and children 2 years of age and older: apply to affected area no more than 3 to 4 times daily ■ children under 2 years of age: ask a doctor</p> <p>Other information ■ store at 20°-25°C (68°-77°F)</p> <p>Inactive ingredients cetyl alcohol, diazolidinyl urea, methylparaben, PEG-2 stearate, PEG-20 stearate, propylene glycol, propylparaben, purified water</p> <p>Questions or comments? 1-800-719-9260</p>
--	--

* This product is not manufactured or distributed by Kenvue Inc., owner of the registered trademark Benadryl®.



ANTI ITCH			
diphenhydramine hydrochloride, zinc acetate cream			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41250-622
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)			
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)			
METHYLPARABEN (UNII: A2I8C7HI9T)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
PROPYLPARABEN (UNII: Z8IX2SC1OH)			
WATER (UNII: 059QF0KO0R)			
DIETHYLENE GLYCOL MONO- AND DIPALMITOSTEARATE (UNII: 94YQ11Y95F)			
PEG-20 STEARATE (UNII: NBX892EA57)			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41250-622-16	2 in 1 CARTON	04/20/2010	06/01/2018
1		28 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:41250-622-64	1 in 1 CARTON	03/17/2010	
2		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 Drug	M017	03/17/2010	

Labeler - Meijer Distribution Inc (006959555)

Revised: 10/2024

Meijer Distribution Inc