

ANTI-ITCH- diphenhydramine hydrochloride and zinc acetate cream

Amerisource Bergen

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.

Anti-Itch Cream

Drug Facts

<i>Active ingredients</i>	<i>Purpose</i>
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

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

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 often 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

- To open: unscrew cap, use pointed end of cap to puncture seal.
- store at 20° to 25°C (68° to 77°F)
- see carton or tube crimp for lot number and expiration date

Inactive ingredients

cetyl alcohol, glyceryl stearate, glyceryl stearate/PEG-100 stearate, methylparaben, propylene glycol, propylparaben and purified water

Questions?

Call **1-866-923-4914**

Distributed By
AmerisourceBergen
1300 Morris Drive
Chesterbrook, PA 19087

PRINCIPAL DISPLAY PANEL - 28.4 g Tube Carton

**GOOD
NEIGHBOR
PHARMACY®**

**Compare to Extra Strength Benadryl®
Itch Stopping Cream* active ingredients***

NDC 24385-210-03

Anti-Itch Cream

Diphenhydramine hydrochloride 2% and Zinc acetate 0.1%

Topical Analgesic • Skin Protectant

NET WT 1 oz (28.4 g)



Compare to Extra Strength Benadryl®
Itch Stopping Cream* active ingredients*

NDC 24385-210-03



Anti-Itch Cream

Diphenhydramine hydrochloride 2% and Zinc acetate 0.1%

Topical Analgesic • Skin Protectant

Relieves pain and itch from insect bites, minor skin irritations
and rashes due to poison ivy, poison oak and poison sumac



NDC 24385-210-03

Anti-Itch Cream

Diphenhydramine hydrochloride 2% and Zinc acetate 0.1%

Topical Analgesic • Skin Protectant



Compare to Extra Strength Benadryl®
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Anti-Itch Cream

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Topical Analgesic • Skin Protectant

NET WT 1 oz (28.4 g)



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Chesterbrook, PA 19087
Visit us at
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Made in Canada

Drug Facts (continued)

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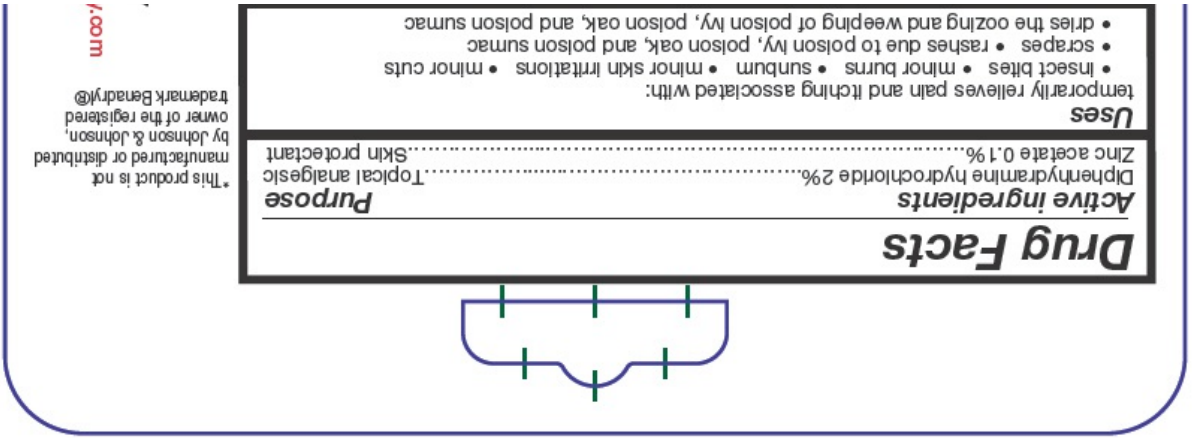
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ANTI-ITCH

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Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:24385-210	
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	
cetyl alcohol (UNII: 936JST6JCN)				
glyceryl monostearate (UNII: 230OU9XXE4)				
methylparaben (UNII: A2I8C7HI9T)				
propylene glycol (UNII: 6DC9Q167V3)				
propylparaben (UNII: Z8IX2SC1OH)				
water (UNII: 059QF0K00R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24385-210-03	1 in 1 CARTON	09/20/2005	
1		28.4 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	09/20/2005	

Labeler - Amerisource Bergen (007914906)

Registrant - Taro Pharmaceuticals U.S.A., Inc (145186370)

Establishment

Name	Address	ID/FEI	Business Operations
Taro Pharmaceuticals Inc.		206263295	MANUFACTURE(24385-210)

Revised: 6/2020

Amerisource Bergen