ANTI-ITCH- diphenhydramine hydrochloride and zinc acetate cream
Amerisource Bergen

Anti-Itch Cream

Drug Facts

Active ingredients	***Purpose***
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)

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

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

GOOD NEIGHBOR PHARMACY®

NDC 24385-210-03

Anti-Itch Cream

Diphenhydramine hydrochloride 2% and Zinc acetate 0.1%

Topical Analgesic • Skin Protectant

LPK-5931-3
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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)

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Drug Facts (continued)

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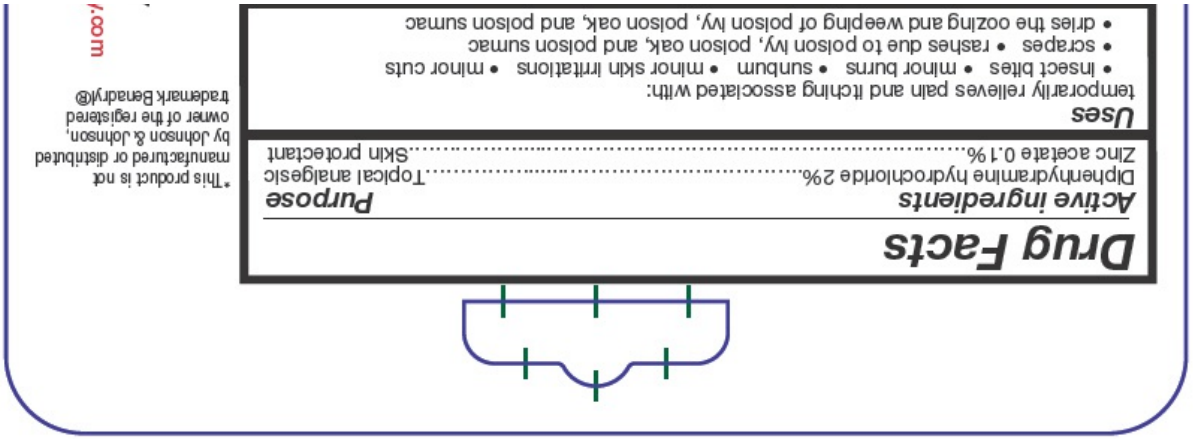
SATISFACTION GUARANTEED
GOOD NEIGHBOR PHARMACY BRAND PRODUCTS

Distributed By
AmerisourceBergen
1300 Morris Drive
Chesterbrook, PA 19087
Visit us at
www.goodneighborpharmacy.com
Made in Canada

NO COPY / NO COLOR
THIS FLAP FOR LOT #
AND EXP DATE PRINT



ANTI-ITCH

diphenhydramine hydrochloride and zinc acetate cream

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: 059QF0KO0R)	

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 Drug	M017	09/20/2005	

Labeler - Amerisource Bergen (007914906)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Pharma Canada Inc.		243339023	manufacture(24385-210)

Revised: 7/2025 Amerisource Bergen