

DIPHENHYDRAMINE HYDROCHLORIDE AND ZINC ACETATE- diphenhydramine hydrochloride and zinc acetate cream
Sun Pharmaceutical Industries, Inc.

Anti-Itch Cream 2% Diphenhydramine Hydrochloride

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:

Taro Pharmaceuticals

U.S.A., Inc.

Hawthorne, NY 10532

PRINCIPAL DISPLAY PANEL - 28.4 g Tube Carton

Extra Strength

Anti-Itch Cream

**Diphenhydramine Hydrochloride 2%
and Zinc Acetate 0.1%**

Topical Analgesic • Skin Protectant

NET WT 1 oz (28.4 g)

NDC 51672-2089-2

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

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

LPK-5891-2
0514-2
M288

Anti-Itch Cream

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Extra Strength

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NET WT 1 oz (28.4 g)

Anti-Itch Cream
Diphenhydramine Hydrochloride 2% and Zinc Acetate 0.1%
Topical Analgesic • Skin Protectant

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Distributed by:
Taro Pharmaceuticals U.S.A., Inc.
Hawthorne, NY 10532

TARO is a registered trademark of Taro Pharmaceuticals U.S.A., Inc.
Made in Canada

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AND EXP DATE PRINT

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This product is not manufactured or distributed by Johnson & Johnson, owner of the registered trademark Benadryl®

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DIPHENHYDRAMINE HYDROCHLORIDE AND ZINC ACETATE

diphenhydramine hydrochloride and zinc acetate cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51672-2089
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)	
PEG-100 STEARATE (UNII: YD01N1999R)	
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:51672-2089-2	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 - Sun Pharmaceutical Industries, Inc. (146974886)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Pharma Canada Inc.		243339023	manufacture(51672-2089)

Revised: 7/2025

Sun Pharmaceutical Industries, Inc.