

LIVE BETTER ANTI-ITCH- diphenhydramine hydrochloride and zinc acetate cream
The Great Atlantic & Pacific Tea Company

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.

Live Better
Anti-Itch

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 ONPOINT, INC.
2 PARAGON DRIVE,
MONTVALE, NJ 07645

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 (28.4g)

NDC 51143-066-02

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



Relieves Pain and Itch from Insect Bites,
Minor Skin Irritations and Rashes due to
Poison Ivy, Poison Oak and Poison Sumac

LPK-7197-0
0113-0
M387

Extra Strength
Anti-Itch Cream

**DIPHENHYDRAMINE HYDROCHLORIDE 2%
AND ZINC ACETATE 0.1%**

■ TOPICAL ANALGESIC ■ SKIN PROTECTANT

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T51



Drug Facts (continued)
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Inactive Ingredients cetyl alcohol, glyceryl stearate, glyceryl stearate/PEG-100 stearate, methylparaben, propylene glycol, propylparaben and purified water.
Questions? Call 1-866-923-4014

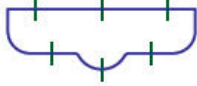
NO COPY / NO COLOR
THIS FLAP FOR LOT #
AND EXP DATE PRINT



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2 PARAGON DRIVE,
MONTVALE, NJ 07645
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Made in Canada

*This product is not
manufactured or distrib-
uted by Johnson & Johnson,
owner of the registered
trademark Benadryl®

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51143-066
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 cation	1 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYL PARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51143-066-02	1 in 1 CARTON		
1		28.4 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part348	09/20/2005	

Labeler - The Great Atlantic & Pacific Tea Company (001367366)**Registrant** - Taro Pharmaceuticals U.S.A., Inc. (145186370)**Establishment**

Name	Address	ID/FEI	Business Operations
Taro Pharmaceuticals Inc.		206263295	MANUFACTURE(51143-066)