EQUALINE ANTI-ITCH- diphenhydramine hydrochloride and zinc acetate cream
Supervalu Inc

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.

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equaline®
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 room temperature
- 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 1-877-932-7948

DISTRIBUTED BY
SUPERVALU INC.
EDEN PRAIRIE,
MN 55344 USA

PRINCIPAL DISPLAY PANEL - 28 g Tube Carton

equaline®

compare to Benadryl®
active ingredients*

extra strength

anti-itch cream
2% diphenhydramine HCl

- topical analgesic/antihistamine
- skin protectant

NET WT 1 OZ (28 g)

equaline®

compare to Benadryl® active ingredients*

extra strength

anti-itch cream

2% diphenhydramine HCl

- topical analgesic/antihistamine
- skin protectant

NET WT 1 OZ (28g)

relieves pain and itch from insect bites, minor skin irritations and rashes from poison ivy, poison oak and poison sumac

equaline®

compare to Benadryl® active ingredients*

extra strength

anti-itch cream

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Drug Facts (continued)

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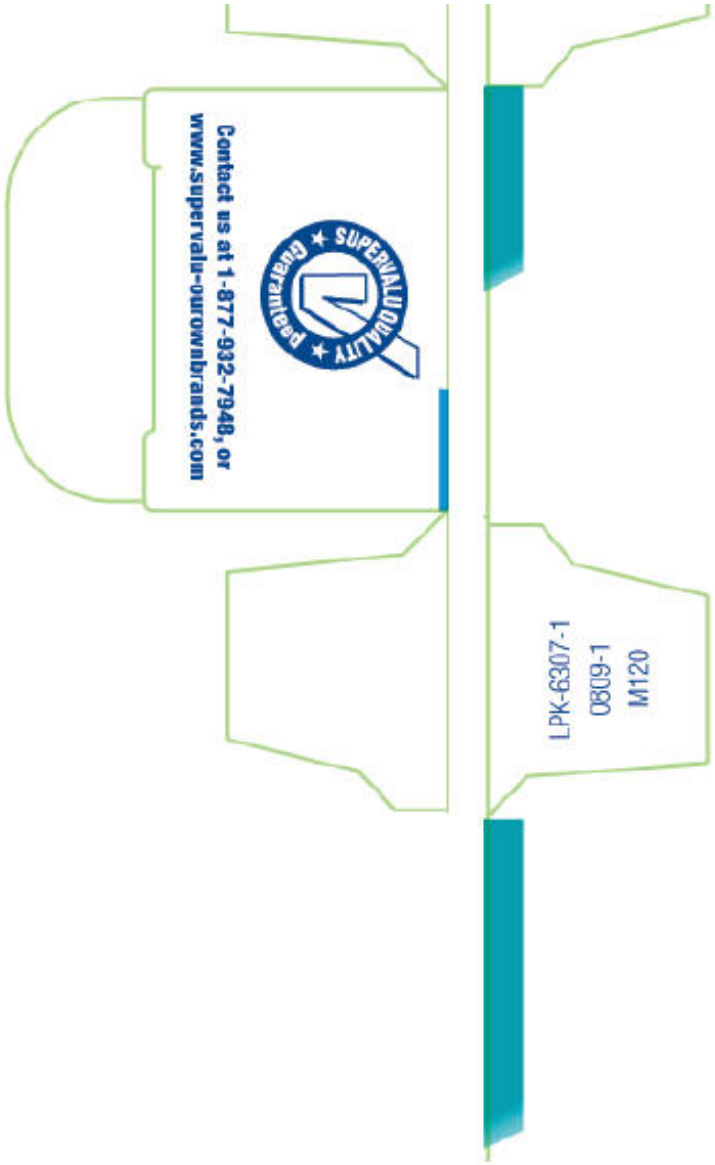

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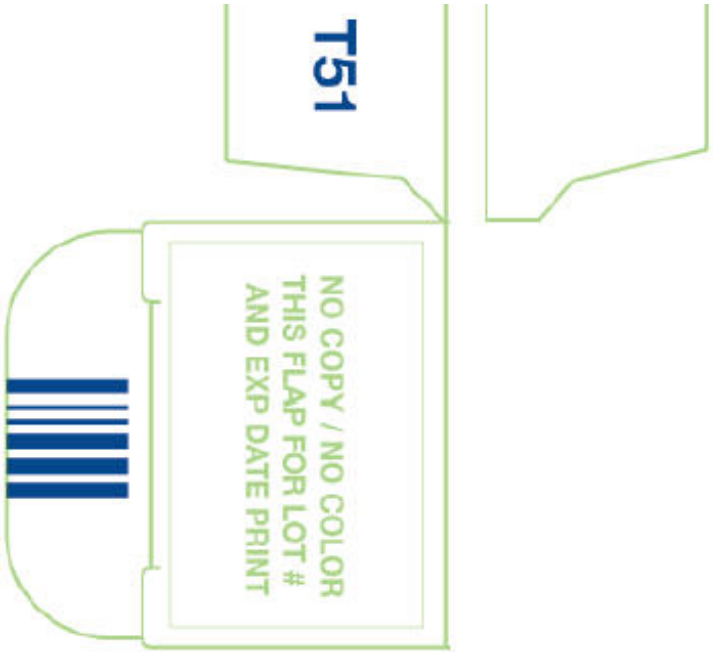
*This product is not manufactured or distributed by Warner-Lambert Consumer Healthcare, owner of the registered trademark Benadryl®

DISTRIBUTED BY
SUPERVALU INC.
EDEN PRAIRIE,
MN 55344 USA
MADE IN CANADA



EQUALINE ANTI-ITCH

diphenhydramine hydrochloride and zinc acetate cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41163-089
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:41163-089-02	1 in 1 CARTON	09/20/2005	
1		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 not final	part348	09/20/2005	

Labeler - Supervalu Inc (006961411)

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

Establishment

Name	Address	ID/FEI	Business Operations
Taro Pharmaceuticals Inc.		206263295	MANUFACTURE(41163-089)

Revised: 2/2019 Supervalu Inc