ANTI-ITCH- diphenhydramine hydrochloride and zinc acetate cream
Target Corporation

Target Up &Up Anti-itch Cream

Drug Fact

Active Ingredients

Diphenhydramine HCL 2%

Zinc Acetate 0.1%

Purpose

Topical analgesic

Skin protectant

Uses

For temporary relief from pain and itching associated with :

- insect bites
- minor burns
- sunburn
- minor skin irritations
- rashes due to poison ivy, poison oak, and poison sumac

Dries the oozing and weeping of poison

- ivy
- oak
- 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 the 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

- conditions worsen or does not improve within 7 days
- symptoms persist for more than 7 days or clear up and occur agian within a few days

Keep this and all drugs out of the reach of children. If swallowed get medical help or contact a Poison Control Center immediately.

Directions

- do not use more than directed
- adults and children 2 years of age and older; apply to affected areas not more than 3 to 4 times daily
- children under 2 years of age , consult a doctor

Other information

Store at controlled room temperature 20°-25°C (68° - 77°F)
Close the cap tightly after use

Inactive ingredients

Cetyl Alcohol, Citric Acid, Diazolldinyl Urea, Glyceral Stearate, Methylparaben, Polyethylene Glycol Monsterate 1000, Sodium Citrate, Water

(Purified)

Questions or comments? 1-800-910-6874

Principal Display Panel

Target Up&Up NDC 11673-865-28

Extra Strength Itch Relief

Diphenhydramine HCL 2% / Zinc Aceate 0.1%

Anti-itch cream

NET WT 1 OZ (28g)

Extra Strength Itch Relief

Diphenhydramine HCL/Topical analgesic
Zinc acetate/Skin protectant

Relieves itches from insect bites and skin irritations

up&up™

NET WT 1 OZ (28 g)

NDC 11673-895-28

Active ingredients Diphenhydramine hydrochloride 2%, Zinc acetate 0.1% ***Purpose*** Topical analgesic, Skin protectant ***Uses*** Temporarily relieves pain and itching associated with: ▪ insect bites ▪ minor burns ▪ sunburn ▪ minor skin irritations ▪ rashes due to poison ivy, poison oak, and poison sumac. Dries the oozing and weeping of poison: ▪ ivy ▪ oak ▪ 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 the 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 this and all drugs out of the reach of children.** If swallowed, get medical help or contact a Poison Control Center immediately. ***Directions*** ▪ do not use more 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: consult a doctor ***Other information*** ▪ Store at controlled room temperature 20°- 25°C (68°- 77°F) ▪ Close the cap tightly after use ***Questions or comments?*** **1-800-910-6874**

245 07 0196 R00 C-002262-01-035

ANTI-ITCH

diphenhydramine hydrochloride and zinc acetate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-895
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)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYETHYLENE GLYCOL 10000 (UNII: H57W405143)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-895-28	1 in 1 CARTON	01/31/2024	
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 Drug	M017	01/31/2024	

Labeler - Target Corporation (006961700)

Registrant - Sheffield Pharmaceuticals LLC (151177797)

Establishment

Name	Address	ID/FEI	Business Operations
Sheffield Pharmaceuticals LLC		151177797	MANUFACTURE(11673-895) , analysis(11673-895)