DERMAMINE EXTRA STRENGTH ITCH STOPPING- diphenhydramine and zinc acetate cream
Natureplex LLC

Dermamine
Extra Strength Itch Stopping Cream

Drug Facts

Active ingredients	***Purposes***
Diphenhydramine Hydrochloride 2%	Topical Analgesic
Zinc Acetate 0.1%	Skin Protectant

Uses

- Temporarily relieves pain and itching associated with:
  - insect bites
  - minor burns
  - sunburn
  - scrapes
  - minor skin irritations
  - minor cuts
  - 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 the reach of children. In case of accidental ingestion, seek medical attention right away 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: ask a doctor

Other information

- store between 15° and 30° C (59° and 86° F)
- close cap tightly after use
- do not use if seal on tube is punctured or missing

Inactive ingredients

DMDM Hydantoin, GMS, isopropyl myristate, light mineral oil, methylparaben, propylparaben, ritacol, purified water

Questions or comments?

1-866-323-0107

PRINCIPAL DISPLAY PANEL - 35 g Tube Box

EXTRA STRENGTH

BLOCKS
THE ITCH-CAUSING
HISTAMINES

Natureplex™

Dermamine
Itch Stopping
Cream
Topical Analgesic Skin Protectant

NDC 67234-023-01

NET WT. 1.25 Oz.(35g)

DERMAMINE EXTRA STRENGTH ITCH STOPPING

diphenhydramine and zinc acetate cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67234-023
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE (UNII: 8GTS82S83M) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE	0.02 mg in 1 g
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	0.001 U in 1 g

Inactive Ingredients

Ingredient Name	Strength
DMDM HYDANTOIN (UNII: BYR0546TOW)	
GLYCERYL 1-STEARATE (UNII: 258491E1RZ)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67234-023-01	1 in 1 BOX	03/01/2014	
1		35 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	03/01/2014	

Labeler - Natureplex LLC (062808196)

Establishment

Name	Address	ID/FEI	Business Operations
Natureplex LLC		062808196	MANUFACTURE(67234-023)