EXTRA STRENGTH ITCH STOPPING- itch stopping cream cream
Universal Distribution Centre LLC

EXTRA STRENGTH ITCH STOPPING CREAM

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

Drug Facts

Active ingredients	**Purpose**
Diphenhydramine hydrochloride 2%	Topical analgesic
Zinc acetate 0.1%	Skin protectant

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 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

- **protect from excessive heat (40°C/104°F)**

Inactive ingredients

cetyl alcohol, diazolidinyl urea, methylparaben, polyethylene glycol monostearate 1000, propylene glycol, propylparaben, purified water

Questions?

Call toll-free 1-800-222-1222

Distributed by:

Universal Distribution Centre LLC

96 Distribution Boulevard, Edison
NJ 08817

PRINCIPAL DISPLAY PANEL - 35.4 g Tube Carton

86073 1.25OZ ITCH STOPPING CREAM - 24

CYAN MAGENTA YELLOW BLACK

Size : 60 X 30 X 40 X 100 X 153 MM

EXTRA STRENGTH

Topical Analgesic/Skin Protectant

ITCH

STOPPING

CREAM

NET WT 1.25 OZ (35.4 g)

EXTRA STRENGTH ITCH STOPPING

itch stopping cream cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52000-071
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
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-071-04	1 in 1 CARTON	01/21/2021	
1	NDC:52000-071-03	35.4 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	M016	01/21/2021	

Labeler - Universal Distribution Centre LLC (019180459)

Registrant - Savvy Care and Cosmetics Pvt. Ltd. (915039748)

Establishment

Name	Address	ID/FEI	Business Operations
Savvy Care and Cosmetics Pvt. Ltd.		915039748	manufacture(52000-071)

Revised: 5/2024 Universal Distribution Centre LLC