

BEST CHOICE ANTI-ITCH CLEAR- zinc acetate and pramoxine hydrochloride lotion
Valu Merchandisers, CO

Best Choice Anti-Itch Clear Lotion

Drug Facts

Active Ingredients

Zinc Acetate 8%

Pramoxine HCl 1%

Purpose

Skin Protectant

External analgesic

Uses

Dries the oozing and weeping, and temporarily relieves pain and itching of poison ivy, oak, and sumac or other skin irritations.

Warnings

For external use only. Use only as directed.

When using this product. Avoid contact with eyes and mucous membranes.

Ask a doctor before using on children 2 years of age.

Stop use and ask a doctor if

condition worsens. Symptoms last for more than 7 days or clear up and occur again within a few days.

Keep out of reach of children.

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Directions

Adults and children 2 yr. of age and older. Shake well before using. Cleanse the skin with soap and water and let dry. Apply to the affected area using cotton or soft cloth, not

more than 3 to 4 times daily as needed for comfort.

Children under 2 yrs. of age. Consult a doctor before use.

Inactive Ingredients

SD Alcohol 38B 2.5%, Camphor, Diazolidinyl Urea, Fragrances, Glycerin, Hydroxypropyl Methycelulose, Methylparaben, Polysorbate 80, Propylene Glycol, Propylparaben and Purified Water.

Other information

Store at room temperature 15-30C (59-86F)

Label

Best Choice
Compare to the Active Ingredients of Caladryl® Clear® Lotion

Anti-Itch Clear

Topical Analgesic • Skin Protectant
Drying Agent • Itch Relief Lotion
6 fl oz (177 mL)

Drug Facts

Active Ingredients
Pramoxine HCl 1%.....External analgesic
Zinc Acetate 0.1%.....Skin protectant

Purpose
External analgesic.....Skin protectant

Indication Dries the oozing and weeping, and temporarily relieves pain and itching of poison ivy, poison oak, and poison sumac or other minor skin irritations.

Warnings
■ For external use only. Use only as directed.
■ Avoid contact with eyes and mucous membranes.
■ Ask a doctor before using on children under 2 years of age.
■ When using this product discontinue use if condition worsens or does not improve or if symptoms persist for more than 7 days, or clear up and occur again within a few days, and consult a doctor.
■ Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Directions
■ Adults and children 2 yrs. of age and older. Shake well before using. Cleanse the skin with soap and water and let dry before each use. Apply lotion to the affected area using cotton or soft cloth, not more than 3 to 4 times daily as needed for comfort.
■ Children under 2 yrs. of age: Consult a doctor before use.

Other information Store at room temperature.

Inactive ingredients SD Alcohol 38B 2.5%, Camphor, Diazolidinyl Urea, Fragrances, Glycerin, Hydroxypropyl Methycelulose, Methylparaben, Polysorbate 80, Propylene Glycol, Propylparaben, Purified Water

*This product is not manufactured or distributed by Johnson & Johnson Consumer Products Company, owner of the registered trademark Caladryl® Clear® Lotion.
R041019RLG

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KANSAS CITY, MO 66106

BEST CHOICE
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BEST CHOICE ANTI-ITCH CLEAR

zinc acetate and pramoxine hydrochloride lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63941-400
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	80 mg in 1 mL
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
CAMPHOR (NATURAL) (UNII: N20HL7Q941)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63941-400-96	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/26/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	03/25/1998	

Labeler - Valu Merchandisers, CO (868703513)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	analysis(63941-400) , manufacture(63941-400) , pack(63941-400) , label(63941-400)

Revised: 12/2023

Valu Merchandisers, CO