PUBLIX CLEAR ANTI-ITCH- zinc acetate and pramoxine hydrochloride lotion Publix

Publix Clear Anti-Itch Lotion

Drug Facts

Active Ingredients

Zinc Acetate 8%

Pramoxine HCI 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 moucous 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 whitin a few days.

Keep out of reach of children.

In case of accidental ingestion, seek profesional 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





ANTI-ITCH LOTION

Skin Protectant

Compare to Active Ingredients in Caladryl® Clear™ Lotion

external analgesic · skin protectant drying agent & itch relief



Drug Facts

Active Ingredients Pramoxine HCI 1% and

Zinc Acetate 0.1%.....External Analgesic/Skin Protectant

Purpose

Uses 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 eye and mucous membranes. ■ Ask a doctor before using on children under 2 years of age.

When using this product

Discontinue use if condition worsens, 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. 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.

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

Other information: Store at room temperature 15° - 30° (59°-86°F).

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Basis of Strength

ZINC CATION

Strength

80 mg

PUBLIX CLEAR ANTI-ITCH

zinc acetate and pramoxine hydrochloride lotion

Product Information

HUMAN OTC DRUG NDC:41415-400 **Product Type** Item Code (Source)

TOPICAL **Route of Administration**

Active Ingredient/Active Moiety Ingredient Name ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)

in 1 mL PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE -**PRAMOXINE** 10 mg UNII:068X84E056) **HYDROCHLORIDE** 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: 60ZP39ZG8H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	

l	Packaging					
	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1 NDC:41415-400-96	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 - Publix (006922009)

Registrant - Pharma Nobis, LLC (118564114)

Establishment			
Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	analysis(41415-400), manufacture(41415-400), pack(41415-400), label(41415-400)

Revised: 12/2023 Publix