# CVS ITCH RELIEF- pramoxine hydrochloride and zinc acetate lotion CVS Pharmacy

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CVS Itch Relief

#### **CVS Itch Relief**

**Drug Facts** 

## **Active Ingredient**

Pramoxine HCl 1%

Zinc Acetate 0.1%

### **Purpose**

External Analgesic

Skin Protectant

#### Use

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

## Warnings

**For external use only.** Use only as directed. Avoid contact with eyes and mucous membranes, Ask doctor before using in children under 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 yrs. 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 Ingredient**

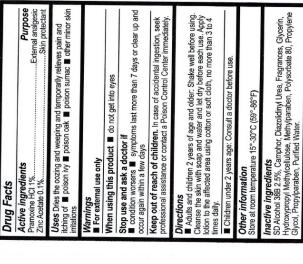
• SD Alcohol 38B 2.5%, Camphor, Diazolidinyl Urea, Fragrances, Glycerin, Hydroxypropyl Methylcellulose, Methylparaben, Polysorbate 80, Propylene Glycol, Propylparaben, and Purified Water.

#### Other Information

• store at room temperature 15  $^{\circ}$  – 30  $^{\circ}$ C (59  $^{\circ}$  – 85  $^{\circ}$ F)

#### PRINCIPAL DISPLAY PANEL







'This product is not manufactured or distributed by Pfizer Consumer Healthcare, owner of the registered trademark Caladryl Clear<sup>®</sup>.

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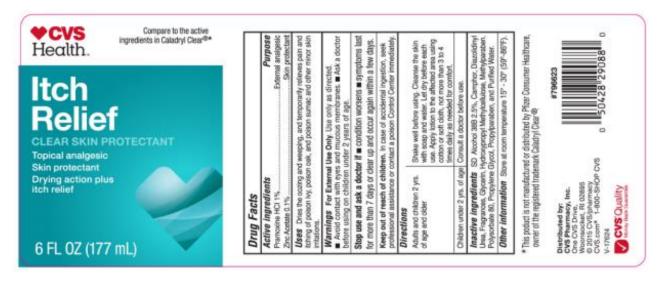
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CVS Health

**Itch Relief** 

#### CLEAR SKIN PROTECTANT

6 FL OZ (177 mL)



## **CVS ITCH RELIEF**

pramoxine hydrochloride and zinc acetate lotion

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Prod	uct	ıntorr	mation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69842-400

**Route of Administration** TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 mL	
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	1 mg in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
ALCOHOL (UNII: 3K9958V90M)			
CAMPHOR (NATURAL) (UNII: N20HL7Q941)			
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)			
GLYCERIN (UNII: PDC6A3C0OX)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
METHYLPARABEN (UNII: A2I8C7HI9T)			
POLYSORBATE 80 (UNII: 60ZP39ZG8H)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
PROPYLPARABEN (UNII: Z8IX2SC10H)			
WATER (UNII: 059QF0KO0R)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842- 400-96	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2008	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	01/01/2008	

## Labeler - CVS Pharmacy (062312574)

## Registrant - Pharma Nobis, LLC (118564114)

Establishment			
Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	analysis(69842-400), manufacture(69842-400), pack(69842-400), label(69842-400)

Revised: 12/2023 CVS Pharmacy