

**CVS CALAMINE PLUS- zinc oxide lotion**  
**CVS Pharmacy**

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**CVS Calamine Plus**

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Drug Facts

**Active Ingredient**

Calamine 8%

**Active Ingredient**

Pramoxine HCl 1%

**Purpose**

Skin Protectant

**Purpose**

External Analgesic

**Use**

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

**Warnings**

For external use only. Use only as directed.

**When using this product avoid contact with eyes.**

**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.

### Other Information

- store at room temperature 15 ° - 30 °C (59 ° - 85 °F)

### Inactive Ingredient

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

### PRINCIPAL DISPLAY PANEL

Calamine  
Plus  
Itch Relief  
6 FL OZ (177 mL)

**CVS Health** Compare to the active ingredients in Caladryl®\*

**Calamine Plus**  
**ITCH RELIEF**  
Topical analgesic  
Skin protectant  
Calamine plus  
Itch reliever

6 FL OZ (177 mL)

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Pramoxine HCl 1%.....External analgesic

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\* This product is not manufactured or distributed by Pfizer Consumer Healthcare, owner of the registered trademark Caladryl®. #109405

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**CVS Pharmacy, Inc.**  
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Money Back Guarantee

## CVS CALAMINE PLUS

zinc oxide lotion

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69842-420
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

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

### Inactive Ingredients

Ingredient Name	Strength
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>CAMPHOR (SYNTHETIC)</b> (UNII: 5TJD82A1ET)	
<b>DIAZOLIDINYL UREA</b> (UNII: H5RIZ3MPW4)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>HYPROMELLOSES</b> (UNII: 3NXW29V3WO)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>WATER</b> (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-420-96	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(69842-420) , manufacture(69842-420) , pack(69842-420) , label(69842-420)