CLEAR ANTI-ITCH- pramoxine hcl, zinc acetate lotion
Consumer Product Partners, LLC

Swan 218.002/218AF
Clear Anti-Itch Lotion

Active ingredients

Pramoxine HCl 1%

Zinc acetate 0.1%

Purpose

External analgesic

Skin protectant

Uses

- for the temporary relief of pain and itching associated with minor skin irritations and rashes due to poison ivy, poison oak or poison sumac
- dries the oozing and weeping of poison: - ivy - oak - sumac

Warnings

For external use only

When using this product

- do not get into eyes

Stop use section and ask a doctor if

- condition worsens
- symptoms last 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

- shake well
- before applying wash affected area of skin

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 - ask a doctor

Other information

store at room temperature (59° - 77°F)

Inactive ingredients

alcohol, benzoic acid, camphor, citric acid, fragrance, glycerin, hydroxypropyl methylcellulose, Lavandula angustifolia (lavender) oil, polysorbate 40, Rosmarinus officinalis (rosemary) leaf oil, sodium citrate, water

Disclaimer

*This product is not manufactured or distributed by Bausch Health US, LLC, distributor of Caladryl ® Clear ® Lotion.

Adverse reaction

Distributed by:

Consumer Product Partners, LLC,

St. Louis, MO 63114

1-888-593-0593

Principal Display Panel

swan ®

Clear Anti-Itch Lotion

External Analgesic/Skin Protectant

Drying action plus itch relief

For relief from pain and itching due to:

- Poison Ivy
- Poison Suman
- Insect Bites
- Poison Oak
- Minor Skin Irritations

Compare to Caladryl ® Clear ® Lotion*

6 FL OZ (177 mL)

CLEAR ANTI-ITCH

pramoxine hcl, zinc acetate lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11344-812
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)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
POLYSORBATE 40 (UNII: STI11B5A2X)	
ROSEMARY OIL (UNII: 8LGU7VM393)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11344-812-30	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/29/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	07/29/2024	

Labeler - Consumer Product Partners, LLC (119091520)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(11344-812)

Revised: 6/2025 Consumer Product Partners, LLC