

CLEAR ANTI ITCH- pramoxine hcl, zinc acetate lotion
United Natural Foods, Inc. dba UNFI

Equaline 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 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 of age - 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 UNFI

PROVIDENCE, RI 02908 USA

855-423-2630

Principal display panel

compare to Caladryl[®] Clear[®] Lotion active ingredients*

NDC 41163-099-30

EQUALINE[®]

clear anti-itch lotion

external analgesic

skin protectant

drying action

plus itch relief

6 FL OZ (177 mL)



clear anti-itch lotion
external analgesic skin protectant

drying action plus itch relief

6 FL OZ (177mL)

compare to Caladryl® Clear® Lotion active ingredients* NDC 41163-099-30

Drug Facts

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L0002471SD

CLEAR ANTI ITCH

pramoxine hcl, zinc acetate lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41163-099
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 (NATURAL) (UNII: N20HL7Q941)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
POLYSORBATE 40 (UNII: ST111B5A2X)	
ROSEMARY OIL (UNII: 8LGU7VM393)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41163-099-30	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/02/2005	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	04/02/2005	

Labeler - United Natural Foods, Inc. dba UNFI (943556183)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091514	manufacture(41163-099)

Revised: 2/2026

United Natural Foods, Inc. dba UNFI