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

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Clear Anti-Itch Lotion
218.002/218AF

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

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

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 (177mL)

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: 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: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 final	part347	04/02/2005	

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

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(41163-099)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(41163-099)

Revised: 5/2023 United Natural Foods, Inc. dba UNFI