

**CLEAR ANTI-ITCH- pramoxine hcl, zinc acetate lotion
Old East Main Co.**

**Rexall 218.002/218AF
Clear Itch Relief 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

Adverse reaction

Visit us at: Rexall.com or call 1-866-4-REXALL

DISTRIBUTED BY OLD EAST MAIN CO.

100 Mission Ridge

Goodlettsville, TN 37072

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A0154

principal display panel

Since 1903

Rexall[®]

Clear Anti-Itch Lotion

External Analgesic • Skin Protectant

For relief from pain and itching due to:

- Poison ivy, oak and sumac
- Insect bites
- Minor skin irritations

6 FL OZ (177 mL)

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Drug Facts

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CLEAR ANTI-ITCH

pramoxine hcl, zinc acetate lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-218
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: 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:55910-218-30	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/30/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	06/30/2011	

Labeler - Old East Main Co. (068331990)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091514	manufacture(55910-218)

Revised: 3/2026

Old East Main Co.