

CALAMINE- calamine, pramoxine hcl lotion
Meijer, Inc.

Meijer 336.003/336AG
Medicated Calamine Lotion

Active ingredients

Calamine 8%

Pramoxine HCl 1%

Purpose

Skin protectant

External analgesic

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

- condition worsens
- symptoms last more than 7 days or clean 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 was 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 - do not use, ask a doctor

Other information

store at room temperature (59°-77°F)

Inactive ingredients

alcohol, benzyl alcohol, camphor, fragrance, hydroxypropyl methylcellulose, Lavandula angustifolia (lavender) oil, phenoxyethanol, polysorbate 80, propylene glycol, Rosmarinus officinalis (rosemary) leaf oil, water, xanthan gum

Disclaimer

*THIS PRODUCT IS NOT MANUFACTURED OR DISTRIBUTED BY BAUSCH HEALTH US, LLC, DISTRIBUTOR OF CALADRYL LOTION CALAMINE PLUS ITCH RELIEVER.

Adverse Reaction

OUR QUALITY GUARANTEE

The Meijer Family

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GRAND RAPIDS, MI 49544

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Principal Panel Display

NDC 79481-0336-1

Compare to Caladryl[®] Lotion Active Ingredients*

meijer[®]

MEDICATED

Calamine

Lotion

External Analgesic

Skin Protectant

For relief from pain and itching due to:

- Poison Ivy

- Poison Sumac
- Insect Bites
- Poison Oak
- Minor Skin Irritation

Empty & Replace Cap

Plastic Bottle

how2recycle.info

6 FL OZ (177 mL)

NDC 79481-0336-1

COMPARE TO CALADRYL LOTION ACTIVE INGREDIENTS*

2

Drying action plus itch relief

Paraben free

Drug Facts

Active ingredients	Purpose
Calamine 8%	Skin protectant
Pramoxine HCl 1%	External analgesic

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 do not use, ask a doctor

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

Inactive ingredients alcohol, benzyl alcohol, camphor, fragrance, hydroxypropyl methylcellulose, Lavandula angustifolia (lavender) oil, phenoxethanol, polysorbate 80, propylene glycol, Rosmarinus officinalis (rosemary) leaf oil, water, xanthan gum

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PID 408021
L0002647SF 0 4 1250 56736 3

CALAMINE

calamine, pramoxine hcl lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79481-0336
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FERRIC OXIDE RED (UNII: 1K09F3G675) (FERRIC OXIDE RED - UNII:1K09F3G675)	FERRIC OXIDE RED	80 mg in 1 mL
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength

ALCOHOL (UNII: 3K9958V90M)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ROSEMARY OIL (UNII: 8LGU7VM393)	
WATER (UNII: 059QF0KO0R)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79481-0336-1	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/17/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	06/17/2009	

Labeler - Meijer, Inc. (006959555)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091514	manufacture(79481-0336)

Revised: 2/2026

Meijer, Inc.