

**CALAMINE - ferric oxide red lotion**  
**Kroger CO**

*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.*

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**Krogers 063.001-063AA**

**Active ingredients**

Calamine 8%  
Zinc oxide 8%

**Purpose**

Skin Protectant

**Use**

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 using
- apply as needed

**Other information**

store at 59° - 86° F

**Inactive ingredients**

bentonite magma, calcium hydroxide, glycerin, purified water

**Questions**

1-800-632-6900

**adverse reactions**

DISTRIBUTED BY THE KROGER CO

CINCINNATI.OHIO, 45202

QUALITY GUARANTEE

[www.kroger.com](http://www.kroger.com)

**principal display panel**

Kroger

FROM OUR FAMILY TO YOURS

SKIN PROTECTANT

calamine lotion

suspension USP

Poison Ivy, Oak and Sumac Drying Lotion

6 FL OZ ( 177 mL)



## CALAMINE

ferric oxide red lotion

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:30142-063
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675) (FERRIC OXIDE RED - UNII:1K09F3G675)	FERRIC OXIDE RED	8 g in 1 mL
<b>ZINC OXIDE</b> (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	8 g in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>BENTONITE</b> (UNII: A3N5ZCN45C)	
<b>CALCIUM HYDROXIDE</b> (UNII: PF5DZW74VN)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>WATER</b> (UNII: 059QF0KO0R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30142-063-30	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/04/2009	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	05/04/2009	

**Labeler** - Kroger CO (006999528)

**Registrant** - Vi Jon, LLC (790752542)

## Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		790752542	manufacture(30142-063)

## Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		088520668	manufacture(30142-063)

Revised: 4/2022

Kroger CO