

**CALAMINE - ferric oxide red lotion**  
**Retail Business Services, LLC**

*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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**Care One 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

**Adverse Reaction Section**

DISTRIBUTED BY FOODHOLD U.S.A., LLC

LANDOVER, MD 20785

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QUALITY GUARANTEED OR YOUR MONEY BACK

**principal display panel**

CAREONE

Calamine

Lotion

CALAMINE TOPICAL

SUSPENSION USP

Skin Protectant

CALAMINE 8%

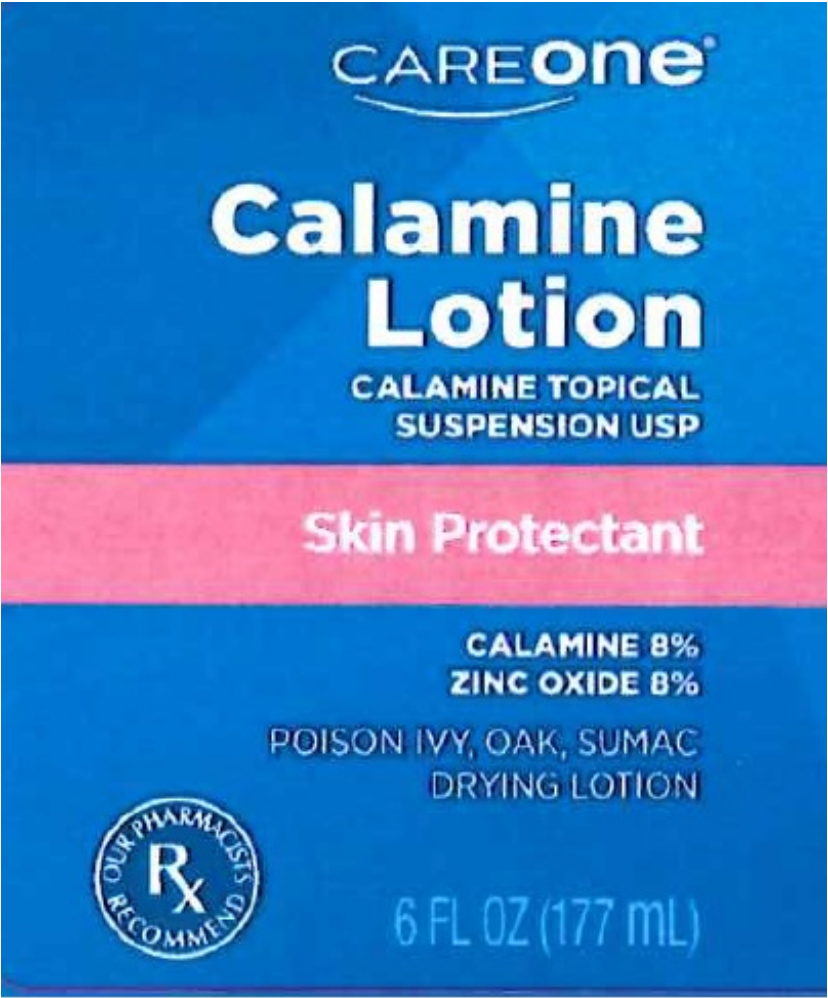
ZINC OXIDE 8%

POISON IVY, OAK, SUMACK

DRYING LOTION

OUR PHARMACISTS RECOMMEND

6 FL OZ (177 mL)



**CALAMINE**

ferric oxide red lotion

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72476-063
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	8 g in 1 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)		ZINC OXIDE	8 mg in 1 mL
Inactive Ingredients			
Ingredient Name			Strength
BENTONITE (UNII: A3N5ZCN45C)			
CALCIUM HYDROXIDE (UNII: PF5DZW74VN)			
GLYCERIN (UNII: PDC6A3C0OX)			
WATER (UNII: 059QF0KO0R)			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72476-063-30	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/06/2013	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	02/06/2013	

**Labeler -** Retail Business Services, LLC (967989935)**Registrant -** Vi Jon, LLC (790752542)**Establishment**

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		088520668	manufacture(72476-063)

**Establishment**

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		790752542	manufacture(72476-063)

Revised: 9/2023

Retail Business Services, LLC