CALAMINE - ferric oxide red lotion Walgreen 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.

Walgreens 063.001/063AA

Active ingredents

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

1-888-593-0593

Disclaimer

Walgreens Pharmacist Recommended
Walgreens Pharmacist Survey Study, November 2016

adverse reactions

DISTRIBUTED BY: WALGREEN CO.

200 WILMOT RD., DEERFIELD, IL

100% SATISFACTION GUARANTEED

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MADE IN U.S.A. WITH U.S. AND FOREIGN COMPENENTS

principal display panel

Walgreens

Calamine

Lotion

CALAMINE TOPICAL

SUSPENSION USP/

SKIN PROTECTANT

• Poison ivy, oak & sumac drying lotion

6 FL OZ (177 mL)



CALAMINE

ferric oxide red lotion

Product	Inform	ation

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-0224
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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: PF5DZ W74VN)				
GLYCERIN (UNII: PDC6A3C0OX)				
WATER (UNII: 059QF0KO0R)				

# Item Code Package Description Marketing Start Date Marketing End Date 1 NDC:0363- 0224-30 177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product 01/05/1996	P	ackaging			
	#	Item Code	Package Description	_	
				01/05/1996	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	01/05/1996	

Labeler - Walgreen Co. (008965063)

Registrant - Vi Jon, LLC (790752542)

Establishme	nt		
Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		790752542	manufacture(0363-0224)

Establishme	ent		
Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		088520668	manufacture(0363-0224)

Revised: 4/2022 Walgreen Co.