CALAMINE - ferric oxide red lotion
Family Dollar

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.

Calamine Lotion
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 59º - 86ºF

Inactive ingredients

bentonite magma, calcium hydroxide, glycerin, purified water

DISTRIBUTED BY: MIDWOOD BRANDS, LLC

500 VOLVO PKWY, CHESAPEAKE,VA 23320 USA

MADE IN THE US WITH US AND FOREIGN COMPONENTS

NOT 100% SATISFIED?

Return package and unused product within 30 days to any Family Dollar store for a refund (with receipt) or exchange.

Principal display panel

FAMILY Wellness

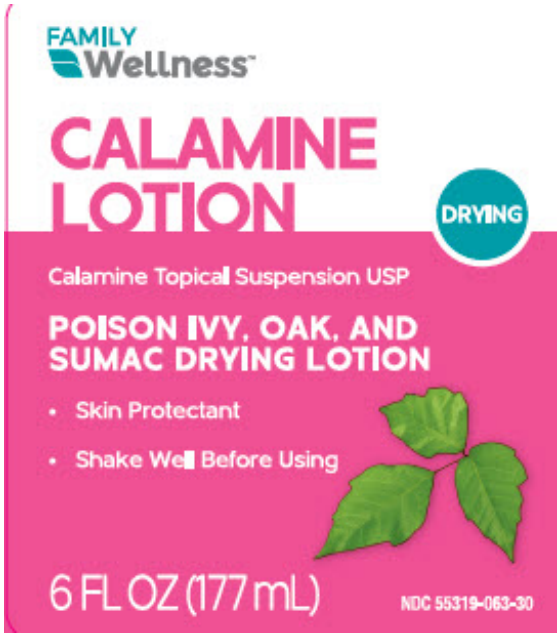
CALMINE LOTION

DRYING

Calamine Topical Suspension USP

POISON IVY, OAK, AND SUMAC DRYING LOTION
- Skin Protectant
- Shake Well Before Using

6 FL OZ (177 mL)



CALAMINE ferric oxide red lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55319-063
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	80 mg in 1 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	80 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:55319-063-30	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1989	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	07/15/1989	

Labeler - Family Dollar (024472631)

Registrant - Vi Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		790752542	manufacture(55319-063)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		088520668	manufacture(55319-063)