CALAMINE- ferric oxide red lotion
Vi Jon, LLC

Swan 063.001/063AA
Calamine Lotion

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 59º - 86ºF

Inactive ingredients

bentonite magma, calcium hydroxide, glycerin, purified water

Adverse reaction

Vi-Jon, LLC

One Swan Drive

Smyrna, TN 37167

Principal Panel Display

NDC 0869-0063-30

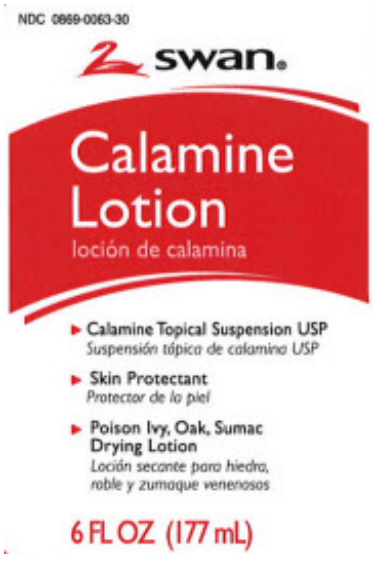
SWAN

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 Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0869-0063
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:0869-0063-30	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1998	
2	NDC:0869-0063-26	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1998	07/01/2017
3	NDC:0869-0063-34	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1998	04/22/2017

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	07/15/1989	

Labeler - Vi Jon, LLC (088520668)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(0869-0063)

Revised: 3/2024 Vi Jon, LLC