CALAMINE PLUS- calamine plus pramoxine hcl aerosol, spray
Chain Drug Marketing Association Inc

Quality Choice Calamine Plus Spray

Active ingredients

Calamine 8%

Pramoxine HCl 1%

Purpose

Skin protectant

External analgesic

Uses

temporarily relieves pain and itching associated with:

- insect bites
- rashes
- minor skin irritations
- minor cuts
- dries the oozing and weeping of poison ivy, poison oak and poison sumac

Warnings

For external use only.

Flammable:

Do not use while smoking or near heat or flame. Do not puncture or incinerate. Contents under pressure. Do not store at temperatures above 120°F. Intentional misuse by deliberately concentrating and inhaling contents can be harmful or fatal.

When using this product

- do not get into eyes
- ask a doctor before using on children under 2 years of age

Stop use and ask a doctor if

- condition worsens
- symptoms last more than 7 days or clear up and occur again in a few days

Keep out of the reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- shake well before using
- adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily
- cleanse the skin with soap and water
- let dry before use
- children under 2 years of age: consult a doctor

Other Information

store between 20° to 25°C (68° to 77°F)

Inactive ingredients

benzyl alcohol, camphor, disteardimonium hectorite, fragrance, hydrated silica, isobutane, oleyl alcohol, SD alcohol 40-B, sorbitan trioleate

Questions?

Call 1-866-964-0939

Principal Display Panel

QC

QUALITY CHOICE

No-Rub

NO-RUB

Calamine

Plus Spray

Calamine 8% / Skin Protectant

Pramoxine HCl 1% / External analgesic

Relieves itching from poison ivy,

oak & sumac, & insect bites

Soothes minor skin irritations & cuts

Shake well before use

NET WT 4.1 OZ. (116 g)

CALAMINE PLUS

calamine plus pramoxine hcl aerosol, spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-242
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	5 mg in 1 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	79.5 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
ISOBUTANE (UNII: BXR49TP611)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
OLEYL ALCOHOL (UNII: 172F2WN8DV)	
SORBITAN TRIOLEATE (UNII: QE6F49RPJ1)	

ALCOHOL (UNII: 3K9958V90M)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-242-41	116 g in 1 CANISTER; Type 0: Not a Combination Product	06/07/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	06/07/2024	

Labeler - Chain Drug Marketing Association Inc (011920774)

Revised: 6/2024 Chain Drug Marketing Association Inc