

**CALAMINE PLUS PRAMOXINE HCL- calamine plus spray aerosol, spray
Foodhold USA**

CareOne Calamine Plus Spray

Active Ingredient

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 temperature 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

- conditions worsens
- symptoms last more than 7 days or clear up and occur again in 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
- adults and children 2 years of age and older: apply as needed to the affected area, not more than 3 or 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**CareOne****NO-RUB CALAMINE SPRAY**

Calamine 8%/ Skin Protectant

Pramoxine HCl 1%/ External Analgesic

TEMPORARILY RELIEVES ITCHING FROM POISON IVY, OAK & SUMAC, AND INSECT BITES

SOOTHES MINOR SKIN IRRITATIONS AND CUTS

Shake well before use

NET WT 4.1 OZ (116g)

CAREone®

NDC XXXXX-XXX-XX

Compare to the active ingredients
in Caladryl® Lotion*

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Drug Facts

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50-07400

CALAMINE PLUS PRAMOXINE HCL

calamine plus spray aerosol, spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41520-184
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	24.16 mg in 116 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	2.72 mg in 116 g
FERRIC OXIDE RED (UNII: 1K09F3G675) (FERRIC OXIDE RED - UNII:1K09F3G675)	FERRIC OXIDE RED	0.345 mg in 116 g

Inactive Ingredients

Ingredient Name	Strength
HYDRATED SILICA (UNII: Y607T4G8P9)	
OLEYL ALCOHOL (UNII: 172F2WN8DV)	
ALCOHOL (UNII: 3K9958V90M)	
SORBITAN TRIOLEATE (UNII: QE6F49RPJ1)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	

ISOBUTANE (UNII: BXR49TP611)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41520-184-41	116 g in 1 CANISTER; Type 0: Not a Combination Product	02/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	02/01/2019	

Labeler - Foodhold USA (809183973)

Revised: 11/2024

Foodhold USA