CALAMINE PLUS PRAMOXINE HCL- calamine plus spray aerosol, spray
Target Corporation

Target 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 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

- 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

No-Rub

Calamine

Spray

Calamine 8% / Skin Protectant

Pramoxine HCL 1% / External

Analgesic

- Soothes minor skin irritations and cuts
- Relieves itching from poison ivy, oak and sumac, and insect bites

Shake Well

Before Use

Temporarily

Relieves

Pain and Itch

NET WT. 4.1 OZ. (116 g)

CALAMINE PLUS PRAMOXINE HCL

calamine plus spray aerosol, spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82442-401
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
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	75 mg in 1 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE -	PRAMOXINE	10 mg

UNII:068X84E056)	HYDROCHLORIDE	in 1 g

Inactive Ingredients

Ingredient Name	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
ISOBUTANE (UNII: BXR49TP611)	
OLEYL ALCOHOL (UNII: 172F2WN8DV)	
SORBITAN TRIOLEATE (UNII: QE6F49RPJ1)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82442-401-01	116 g in 1 CAN; Type 0: Not a Combination Product	05/29/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/29/2025	

Labeler - Target Corporation (006961700)

Revised: 5/2025 Target Corporation