FAMILY CARE MAXIMUM STRENGTH MEDICATED ANTI ITCH- menthol cream
United Exchange Corp.

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.

Family Care Anti-Itch + Pain Relief Cream 0.5oz (10497)

Active Ingredients Purpose

Menthol 1%.....Anti-itch/Topical analgesic

Pramoxine HCl 1%.....Anti-itch/Topical analgesic

Uses
- for temporary relief of pain and itching associated with:
- minor skin irritations
- minor cuts
- minor burns
- minor sunburns
- scrapes
- insect bites
- rashes due to poison ivy, oak, or sumac

Warnings For external use only.

Do not use
- on deep or puncture wounds
- animals bites
- serious burns
- large areas of the body

When using this product, avoid contact with eyes or nose.

Stop use and ask a doctor if
- condition worsens, symptoms persist for more than 7 days or clear up and occur again within a few days
- if redness, irritation, swelling or pain persists or increases

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. If swallowed, get medical help or contact a Poison control Center (1-800-222-1222) right away.

Directions
- not for prolonged use
- adults and children 2 years of age and older; apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: ask a doctor.

Other information
- store at room temperature 20-25°C (68-77°F)

Inactive ingredients aloe vera leaf, carbomer 940, glycerin, glyceryl monostearate, isopropyl myristate, methyl salicylate, methylparaben, mineral oil, polysorbate 60, propylparaben, purified water, sodium hydroxide, stearyl alcohol, tocopheryl acetate

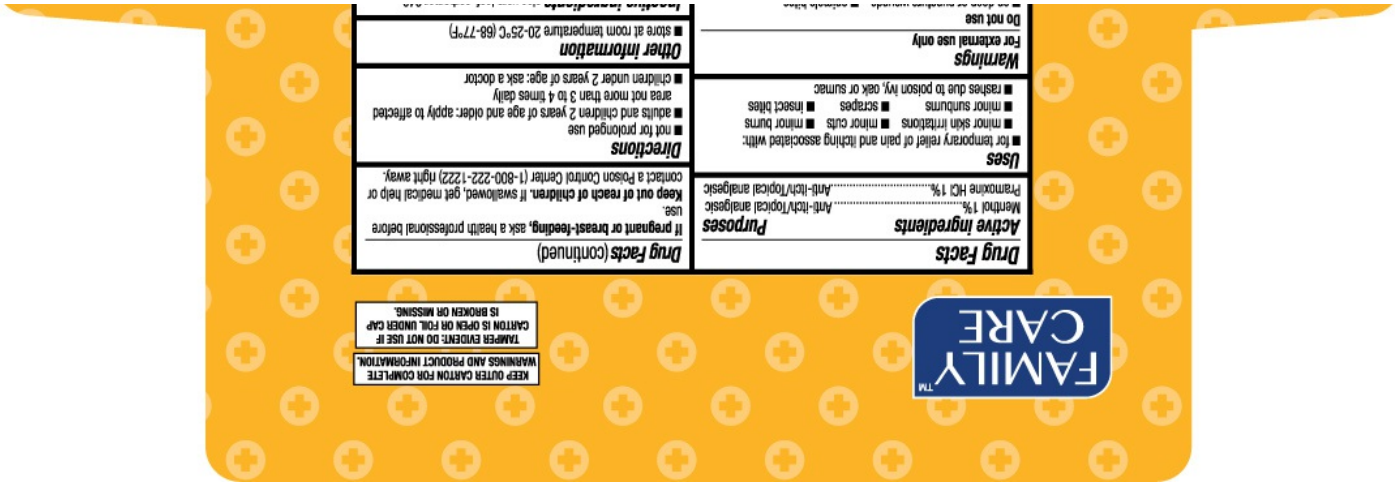
Questions or comments?

Call 1-888-645-8204 Monday-Friday 9AM-5PM (PST)

Distributed By: UNITED EXCHANGE CORPORATION

Cypress, CA 90630 USA | 1-888-645-8204

Made in China



FAMILY CARE MAXIMUM STRENGTH MEDICATED ANTI ITCH

menthol cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65923-104
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	10 mg in 1 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER HOMOPOLYMER TYPE C (UNII: 4Q93RCW27E)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
MINERAL OIL (UNII: T5L8T28FGP)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-104-14	1 in 1 CARTON	10/30/2018	
1		14 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part346	11/24/2014	

Labeler - United Exchange Corp. (840130579)

Revised: 3/2022 United Exchange Corp.