

**QC MEDICATED ANTI ITCH- pramoxine hydrochloride, menthol cream**  
**Chain Drug Marketing Association**

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**QC Medicated Anti Itch Cream**

**Active Ingredient**

Menthol 1%

Pramoxine hydrochloride 1%

**Uses**

for the temporary relief of pain and itching due to: minor burns, sunburn, minor cuts, scrapes, insect bites, minor skin irritations ,minor rashes due to poison ivy, poison oak, or poison sumac

**Purpose**

Topical anagesic

**Warnings**

**For external use only**

**When using this product**

do not get into eyes

**Keep out of reach of children**

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

**Other Information**

store at controlled room temperature

**Directions**

adults and children 2 years and older: apply to affected area not more than 3 to 4 times daily

children under 2 years: consult a doctor

**Dosage**

Adults and Children 2 years and older: Apply to affected are not more than 3 to 4 times daily.

## **Inactive Ingredients**

aloe barbadensis (aloe vera) leaf juice, diazolidinyl urea, edetate disodium, eucalyptus oil, methylparaben, methyl salicylate, mineral oil, PPG-1 trideceth-6, propylene glycol, propylparaben, purified water, sodium acrylates copolymer, steareth-21, stearyl alcohol, tocopheryl acetate, trolamine, white petrolatum

## **Package principal panel**

PMS 200 C  
CMYK



\*Compare to the Active Ingredients  
in Gold Bond® Anti-Itch Cream

# Medicated Anti-Itch Cream

Topical Analgesic with Aloe & Vitamin E

Maximum Strength Pain & Itch Relief  
Steroid Free | Hydrocortisone Free

## Relieves Pain & Itch Fast

Insect  
Bites

Minor Cuts  
& Scrapes

Poison Ivy,  
Oak, Sumac

Sunburn

Skin  
Irritation

NDC 63868-959-01



\*Compare to the Active Ingredients  
in Gold Bond® Anti-Itch Cream

# Medicated Anti-Itch Cream

Topical Analgesic with Aloe & Vitamin E



1 oz (28 g) Net Wt.

QC  
Quality Choice  
Medicated  
Anti-Itch Cream  
Topical Analgesic  
with Aloe & Vitamin E

Lot #

Exp

\*This product is not manufactured or distributed by Chattem, Inc.,  
owner of the registered trademark Gold Bond®.



Distributed by C.D.M.A., Inc.®  
43157 W 9 Mile Rd  
Novi, MI 48375  
www.qualitychoice.com  
Questions: 800-935-2362



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Made in the U.S.A.

05-265-51V3.0

**Drug Facts**

**Active ingredients**  
Menthol 1%  
Pramoxine hydrochloride 1%  
Topical analgesic

**Purposes**  
Topical analgesic

**Uses**  
• for the temporary relief of pain and itching due to:  
• minor burns • sunburn • minor cuts • scrapes • insect bites  
• minor skin irritations • minor rashes due to poison ivy, poison oak, or poison sumac

**Warnings**  
For external use only  
When using this product do not get into eyes  
Stop use and ask a doctor if • condition gets worse • symptoms last for 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**  
• adults and children 2 years and older: apply to affected area not more than 3 to 4 times daily  
• children under 2 years: consult a doctor

**Other information** • store at controlled room temperature • to report serious side effects call (515) 276-1586

**Inactive ingredients** aloe barbadensis (aloe vera) leaf juice, diazolidinyl urea, edetate disodium, eucalyptus oil, methylparaben, methyl salicylate, propylene glycol, propylparaben, purified water, sodium polyacrylate, steareth-21, stearyl alcohol, tocopheryl acetate, tolimaine, white petrolatum

Important: keep this carton —  
it has complete information.

pramoxine hydrochloride, menthol cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63868-959
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>MENTHOL</b> (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 g in 100 g
<b>PRAMOXINE HYDROCHLORIDE</b> (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	1 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>DIAZOLIDINYL UREA</b> (UNII: H5RIZ3MPW4)	
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)	
<b>STEARETH-21</b> (UNII: 53J3F32P58)	
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>SODIUM POLYACRYLATE (8000 MW)</b> (UNII: 285CYO341L)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>METHYL SALICYLATE</b> (UNII: LAV5U5022Y)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>STEARYL ALCOHOL</b> (UNII: 2KR89I4H1Y)	
<b>EUCALYPTUS OIL</b> (UNII: 2R04ONI662)	
<b>.ALPHA.-TOCOPHEROL ACETATE, DL-</b> (UNII: WR1WPI7EW8)	

### Product Characteristics

<b>Color</b>	white	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-959-01	1 in 1 CARTON	12/07/2020	
1		28 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 Drug	M017	12/07/2020	

**Labeler** - Chain Drug Marketing Association (011920774)

**Registrant** - Weeks & Leo Co., Inc. (005290028)

## Establishment

Name	Address	ID/FEI	Business Operations
WEEKS & LEO COMPANY, INC.		005290028	manufacture(63868-959)

Revised: 12/2024

Chain Drug Marketing Association