

**GOLD BOND ANTI ITCH- menthol and pramoxine hydrochloride cream
Chattem, Inc.**

Gold Bond Anti Itch

Drug Facts

Active ingredients

Menthol 1%

Pramoxine hydrochloride 1%

Purpose

Anti-itch, Pain relief

Uses

for temporary relief of pain and itching associated with:

- minor skin irritations
- minor cuts
- minor burns
- minor sunburns
- rashes due to poison ivy, poison oak or poison sumac
- scrapes
- insect bites

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

- do not get into eyes or nose
- not for prolonged use

Stop use and ask a doctor if

- condition worsens
- symptoms last more than 7 days or clear up and occur again within a few days
- if redness, irritation, swelling or pain persists or increases

Keep out of reach of children.

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

Directions

adults and children 2 years and older: apply to affected area up to 3 or 4 times daily

children under 2 years: consult a doctor

Inactive ingredients

water, petrolatum, propylene glycol, stearyl alcohol, aloe barbadensis leaf juice, triethanolamine, sodium acrylates copolymer, steareth-21, mineral oil, steareth-2, tocopheryl acetate, thymol, eucalyptol, methyl salicylate, PPG-1 trideceth-6, diazolidinyl urea, disodium EDTA, , iodopropynyl butylcarbamate

PRINCIPAL DISPLAY PANEL

RAPID RELIEF GOLD BOND®
Net wt 1 oz (28 g)
ANTI-ITCH CREAM



GOLD BOND ANTI ITCH
menthol and pramoxine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41167-0501
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Route of Administration TOPICAL**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.01 g in 1 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	0.01 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PETROLATUM (UNII: 4T6H12BN9U)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TROLAMINE (UNII: 9O3K93S3TK)	
STEARETH-21 (UNII: 53J3F32P58)	
MINERAL OIL (UNII: T5L8T28FGP)	
STEARETH-2 (UNII: V56DFE46J5)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
THYMOL (UNII: 3J50XA376E)	
EUCALYPTOL (UNII: RV6J6604TK)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
PPG-1 TRIDECETH-6 (UNII: 1K7417JX6Q)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41167-0501-0	1 in 1 CARTON	04/01/1996	
1		28 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:41167-0501-1	1 in 1 CARTON	04/01/1996	02/27/2019
2		35 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:41167-0501-2	1 in 1 CARTON	04/01/1996	02/28/2019
3		56 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
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Category	Citation	Date	Date
OTC Monograph Drug	M017	04/01/1996	

Labeler - Chattem, Inc. (003336013)

Revised: 11/2023

Chattem, Inc.