GOLD BOND INTENSIVE RELIEF ANTI-ITCH- pramoxine hydrochloride, menthol lotion
Chattem, Inc.

Gold Bond Intensive Relief Anti-Itch

GOLD BOND® Medicated

Intensive Relief Anti-Itch Lotion

Drug Facts

Active ingredients

Menthol 0.5%
Pramoxine hydrochloride 1%

Purpose

Anti-itch
Anti-itch

Use

for temporary relief of pain and itching associated with:

■ minor burns ■ sunburn ■ minor cuts ■ scrapes ■ insect bites ■ minor skin irritations

Warnings

For external use only

When using this product

■ avoid contact with eyes

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

Keep out of reach of children.

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

Directions

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

children under 2 years of age: consult a doctor

Inactive ingredients

water, glycerin, dimethicone, cetyl alcohol, petrolatum, distearyldimonium chloride, aloe barbadensis leaf juice, glyceryl stearate, stearyl alcohol, cetearyl alcohol, behentrimonium methosulfate, steareth-21, propylene glycol, polysorbate 60, stearamidopropyl PG-dimonium chloride phosphate, diazolidinyl urea, butylene glycol, tocopheryl acetate, potassium hydroxide, methylparaben, panthenol, ethoxydiglycol, EDTA, avena sativa (oat) kernel extract

PRINCIPAL DISPLAY PANEL

GOLD BOND
ANTI-ITCH LOTION
INTENSIVE RELIEF
Net wt 5.5 oz (155 g)

GOLD BOND INTENSIVE RELIEF ANTI-ITCH

pramoxine hydrochloride, menthol lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41167-0507
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	1 g in 100 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
PETROLATUM (UNII: 4T6H12BN9U)	
DISTEARYLDIMONIUM CHLORIDE (UNII: OM9573ZX3X)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
BEHENTRIMONIUM METHOSULFATE (UNII: 5SHP745C61)	
STEARETH-21 (UNII: 53J3F32P58)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
STEARAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: W6000VEI5Y)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PANTHENOL (UNII: WV9CM0O67Z)	
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A1I8X02B)	
EDETIC ACID (UNII: 9G34HU7RV0)	
OAT (UNII: Z6J799EAJK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41167-0507-0	155 g in 1 BOTTLE; Type 0: Not a Combination Product	04/26/2016	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC Monograph Drug	M017	04/26/2016	

Labeler - Chattem, Inc. (003336013)

Revised: 10/2023 Chattem, Inc.