

**GOLD BOND MEDICATED PAIN AND ITCH RELIEF- lidocaine
hydrochloride cream
Chattem, Inc.**

Gold Bond Medicated Pain and Itch Relief

**Gold Bond® Medicated
Pain & Itch Relief Cream with Lidocaine**

Drug Facts

Active ingredient

Lidocaine HCl 4%

Purpose

Topical anesthetic

Use

for temporary relief of the pain and itching associated with:

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

Warnings

For external use only

Do not use

- in large quantities, particularly over raw surfaces of blistered area
- on deep or puncture wounds

When using this product

- use only as directed. Read and follow all directions and warnings on this carton.
- 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 right away.

Directions

adults and children 12 years of age and older:

- apply a thin layer to affected area not more than 3 to 4 times daily

children under 12 years of age: consult a doctor

Inactive ingredients

acrylates/C10-30 alkyl acrylate crosspolymer, aloe barbadensis leaf juice, aminomethyl propanol, C30-45 alkyl cetearyl dimethicone crosspolymer, caprylyl methicone, cetearyl alcohol, ceteth-20 phosphate, dicetyl phosphate, dimethicone, disodium EDTA, ethylhexylglycerin, glyceryl stearate, methylparaben, SD alcohol 40 (15%), steareth-21, water (309-179)

Child resistant packaging. Close cap tightly between uses.

PRINCIPAL DISPLAY PANEL

GoldBond
Pain & Itch Relief Cream
Net Wt 1.75 oz (49g)

PAIN & ITCH FORMULA

- Starts Working on Contact
- Minor Burns & Cuts
- Minor Skin Irritations
- Insect Bites
- Sunburn
- Steroid Free



MULTI-SYMPTOM

WITH **LIDOCAINE** MAXIMUM STRENGTH
4%

GOLD BOND



Net wt 1.75 oz (49 g)

PAIN & ITCH RELIEF CREAM

GOLD BOND
4% LIDOCAINE



NEVER USE IF CONTAINER IS DAMAGED OR IF SEAL IS BROKEN

SANOFI
 Distributed by: **Sanofi**, Inc., 1 Sanofi Square
 P.O. Box 2378, Channahon, IL 61615-0237
 www.goldbond.com 815-331-0100

Child resistant packaging. Close caps tightly between uses.

Drug Facts
Active ingredient Lidocaine HCl 4%
Purpose Topical anesthetic
Use 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
Do not use ■ in large quantities, particularly over the surface or thinned areas ■ on deep or puncture wounds ■ when using this product ■ use only as directed. Read and follow directions and warnings on the carton. ■ 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
Directions Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately. adults and children 12 years of age and older: apply a thin layer to affected areas not more than 3 to 4 times daily children under 12 years of age: consult a doctor
Inactive ingredients acrylates C10-13 alkyl acrylate crosspolymer, acryol denat (15%), iso butyl alcohol, methyl-20 propylate, dicyl propylate, dimethylsiloxane, decadum EDTA, ethylhexylglycerin, glyceryl stearate, methylparaben, stearyl-21, water

GOLD BOND

GOLD BOND

GOLD BOND® Pain & Itch Relief Cream
 with Lidocaine Numbs Away Itch and Pain.
 This Steroid Free, non-irritating formula
 contains Maximum Strength Lidocaine (4%)
 and starts working on contact.

GOLD BOND
4% LIDOCAINE

GOLD BOND MEDICATED PAIN AND ITCH RELIEF

lidocaine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41167-0505
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETETH-20 PHOSPHATE (UNII: 921FTA1500)	
DIHEXADECYL PHOSPHATE (UNII: 2V6E5WN99N)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
ALCOHOL (UNII: 3K9958V90M)	
STEARETH-21 (UNII: 53J3F32P58)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41167-0505-0	1 in 1 CARTON	01/15/2016	
1		49 g in 1 BOTTLE, PLASTIC; 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	01/15/2016	

Labeler - Chattem, Inc. (003336013)

