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

Gold Bond Medicated Pain and Itch Relief - Reformulated

Gold Bond® Medicated

Pain & Itch Relief Cream with Lidocaine

Drug Facts

Active ingredient

Purpose

Lidocaine HCl 4%.....Topical anesthetic

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 raw surfaces or blistered areas ■ on deep or puncture wounds

When using this product

■ use only as directed. Read and follow all directions and warnings on this label.

■ 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 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, alcohol denat. (15%), 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, steareth-21, water

Close cap tightly between uses.

PRINCIPAL DISPLAY PANEL

GOLD BOND
Maximum Strength
Pain and Itch Relief Cream
Net wt.1.75 oz (49 g)

GOLD BOND MEDICATED PAIN AND ITCH RELIEF

lidocaine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41167-0529
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
ALCOHOL (UNII: 3K9958V90M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
C30-45 ALKYL CETEARYL DIMETHICONE CROSSPOLYMER (UNII: 4ZK9VP326R)	
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)	
STEARETH-21 (UNII: 53J3F32P58)	
WATER (UNII: 059QF0KO0R)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41167-0529-0	1 in 1 CARTON	03/01/2023	
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	03/01/2023	

Labeler - Chattem, Inc. (003336013)

Revised: 1/2026 Chattem, Inc.