

LIDOCAINE- lidocaine gel

Bellus Medical, LLC

4% Lidocaine Gel

Lidocaine 4%

Water, Ethyl Alcohol, Glycerin, Propylene Glycol, Xanthan Gum, Phenoxyethanol, Caprylyl Glycol, Aloe Barbadensis (Aloe Vera) Leaf Extract, Sorbic Acid, Tetrasodium EDTA.

USES: For the temporary relief of discomfort and pain associated with

- Minor burns and skin irritations
- Minor cuts and scrapes
- Itching

DIRECTIONS: Adults and Children 12 years of age and older apply to affected area not more than 3-4 times daily.

WARNINGS: For external use only. Avoid contact with the eyes.

DO NOT USE: In large quantities particularly over raw sources or blistered areas.

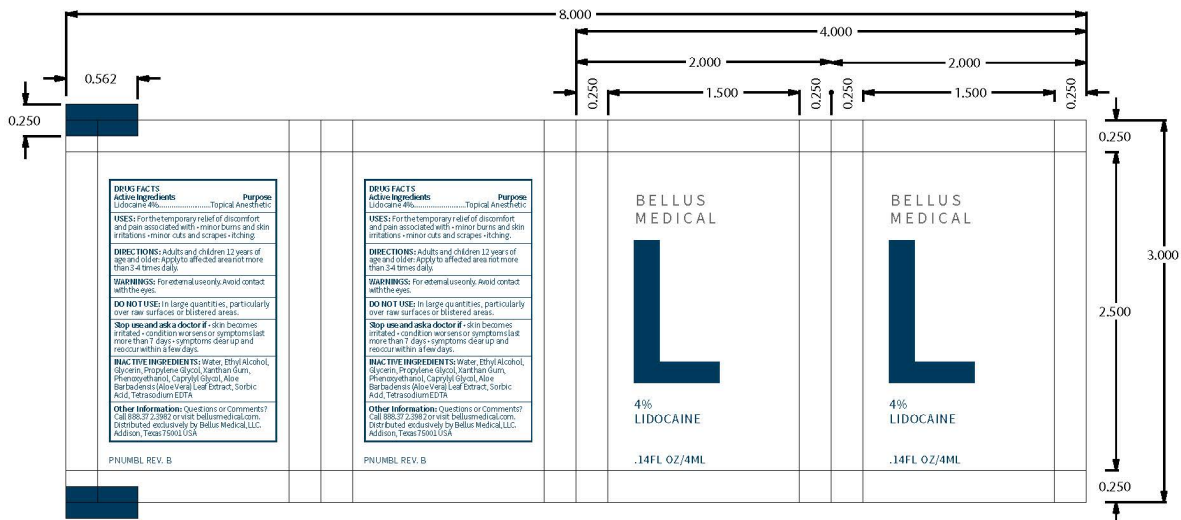
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Topical Anesthetic

Keep put of reach of children.



lidocaine gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71888-102
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SORBIC ACID (UNII: X045WJ989B)	
EDETATE SODIUM (UNII: MP1J8420LU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71888-102-02	12 in 1 BOX	06/01/2017	
1	NDC:71888-102-01	4 mL in 1 PACKET; 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	06/01/2017	

Labeler - Bellus Medical, LLC (005677967)

Registrant - Bellus Medical, LLC (005677967)