DIABETIC FIRST AID GEL- benzalkonium chloride, lidocaine hydrochloride gel Lavior Pharma Inc.

Active Ingredients

Benzalkonium Chloride 0.13%Topical Antiseptic Lidocaine Hydrochloride 0.5%Topical Analgesic

Purpose

Topical Antiseptic Topical Analgesic

Uses

First aid to help prevent infection in minor:

■cuts **■**scrapes **■**burns

Temporarily relieves pain and itching associated with:

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

Warnings

For external use only

Do not use

- in the eyes
- over large areas of the body
- in large quantities, particularly over raw surfaces or blistered areas

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- condition or symptoms get worse or last more than 1 week
- symptoms 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:

- clean the affected area
- apply a small amount of this product on the area 1 to 3 times daily
- may be covered with a sterile bandage
- if bandaged, let dry first

Children under 12 years of age: consult a doctor

Other information

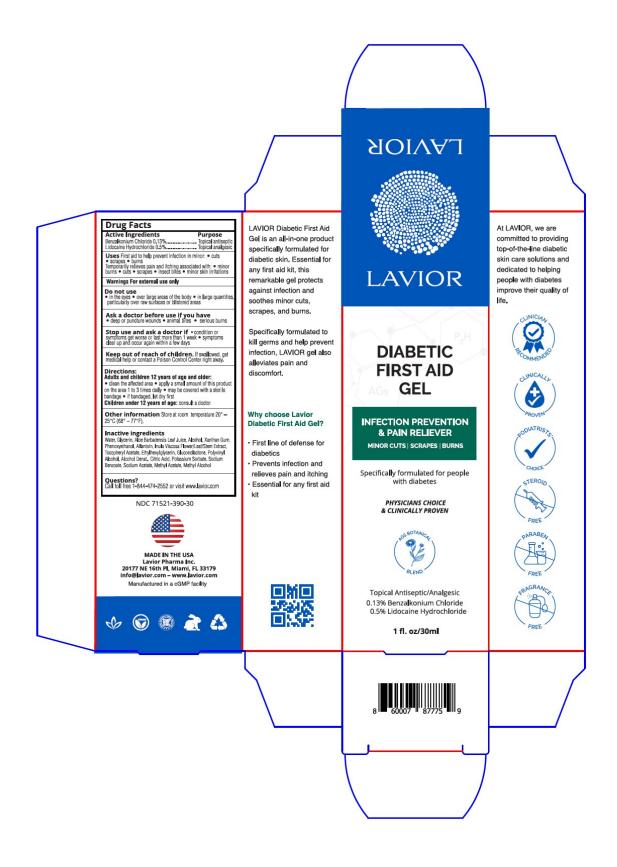
Store at room temperature

Inactive Ingredients

Water, Glycerin, Inula Viscosa Flower/Leaf/Stem Extract, Aloe Barbadensis Leaf Juice, Xanthan Gum, Phenoxyethanol, Allantoin, Tocopheryl Acetate, Ethylhexylglycerin, Gluconolactone, Polyvinyl Alcohol, Alcohol, Citric Acid, Sodium Benzoate, Potassium Sorbate

Questions?

Call (305) 627-3680 or visit www.lavior.com



DIABETIC FIRST AID GEL

benzalkonium chloride, lidocaine hydrochloride gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71521-390

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Route	Ot A	Δdmin	ictrai	ion

TOPICAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	0.5 g in 100 g		
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g		

Inactive Ingredients	
Ingredient Name	Strength
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ALCOHOL (UNII: 3K9958V90M)	
GLUCONOLACTONE (UNII: WQ29KQ9POT)	
GLYCERIN (UNII: PDC6A3C0OX)	
XANTHAN GUM (UNII: TTV12P4NEE)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
ALLANTOIN (UNII: 344S277G0Z)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
DITTRICHIA VISCOSA WHOLE (UNII: 3SYW69FH88)	

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:71521- 390-50	50 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/31/2021		
2	NDC:71521- 390-15	15 g in 1 TUBE; Type 0: Not a Combination Product	03/17/2022		
3	NDC:71521- 390-30	30 g in 1 TUBE; Type 0: Not a Combination Product	04/01/2023		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M003	03/31/2021		

Labeler - Lavior Pharma Inc. (080685327)

Revised: 10/2024 Lavior Pharma Inc.