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

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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

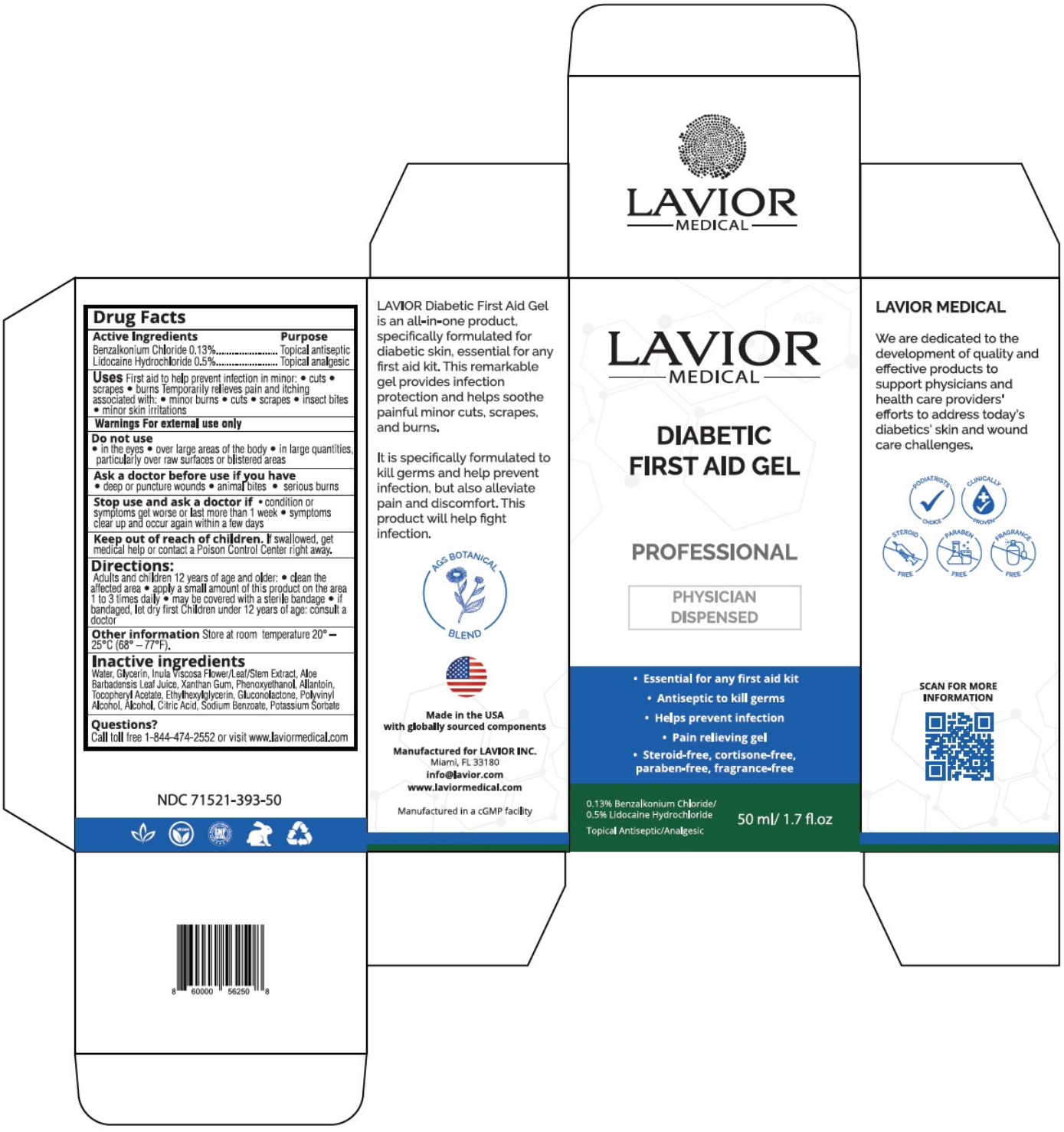
Store at room temperature

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 toll free 1-844-474-2552 or visit www.lavior.com



DIABETIC FIRST AID GEL PROFESSIONAL

benzalkonium chloride, lidocaine hydrochloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71521-393
Route of Administration	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

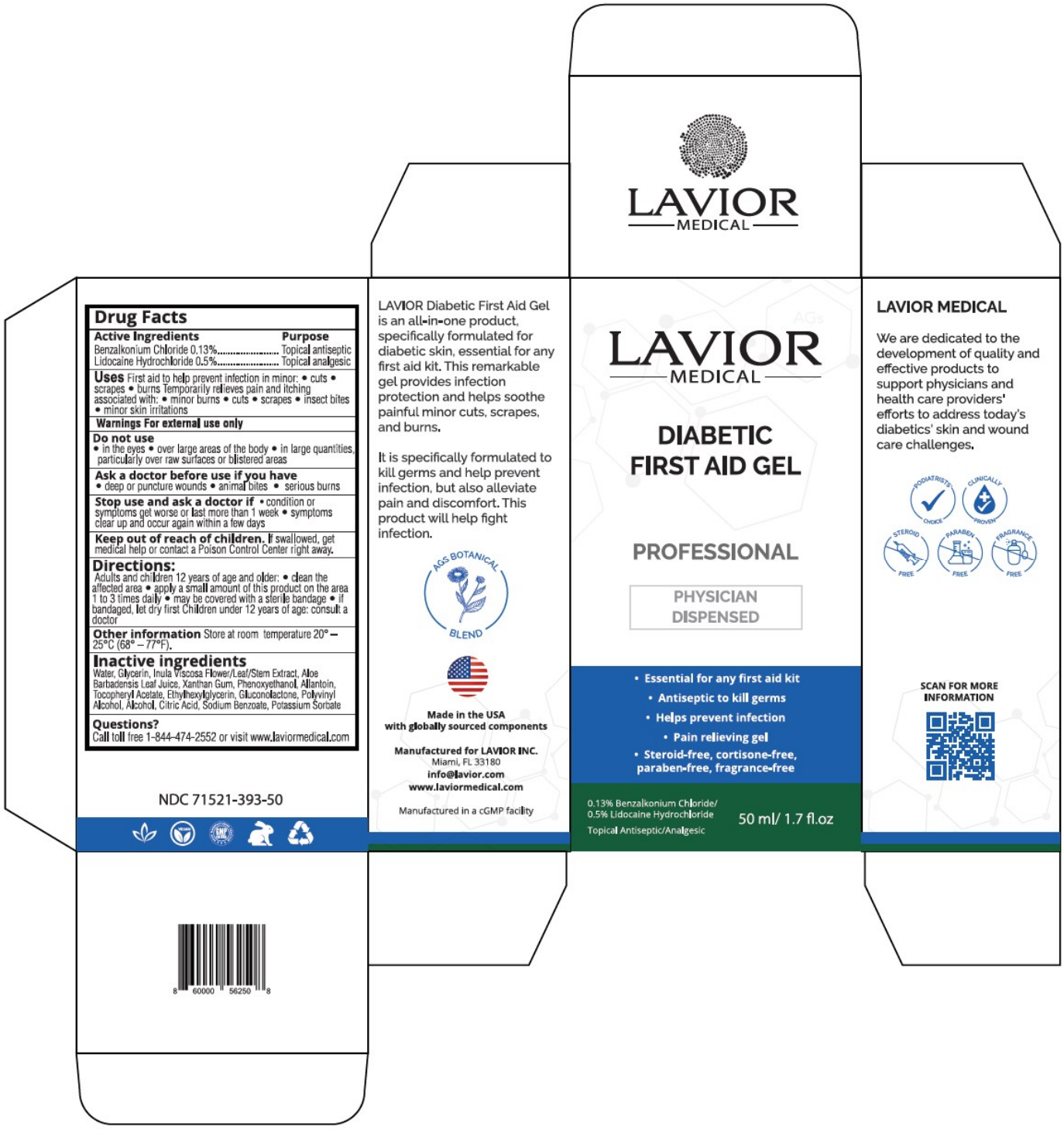
Ingredient Name	Strength
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ALCOHOL (UNII: 3K9958V90M)	
GLUCONOLACTONE (UNII: WQ29KQ9POT)	
GLYCERIN (UNII: PDC6A3C0OX)	
XANTHAN GUM (UNII: TTV12P4NEE)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
CITRIC ACID MONOHYDRATE (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)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71521-393-50	50 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/31/2021	



Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/31/2021	

Labeler - Lavior Inc. (080685327)

Revised: 5/2021 Lavior Inc.