ANTIBACTERIAL LIDOCAINE WOUND GEL- benzalkonium chloride, lidocaine hydrochloride gel
ASO LLC

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.

Antibacterial Lidocaine Wound Gel

Active ingredient (in each gram)

Benzalkonium Chloride 0.13%
Lidocaine Hydrochloride 2.00%

Purpose

first aid antiseptic
external analgesic

Uses

- First aid to help prevent infection in minor cuts, scrapes and burns
- For the temporary relief of pain associated with minor burns

Warnings

For external use only

Do not use

- in the eyes
- over large areas of the body, particularly over raw surfaces or blistered areas
- longer than a week unless directed by a doctor

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- the condition persists more than 7 days or gets worse
- condition clears up and occurs again within a few days

Keep out of reach of children.

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Directions

- clean the affected area
- adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: consult a doctor
- may be covered with a sterile bandage
- if bandaged, let dry first

Other information

Store at room temperature

Inactive ingredients

caprylyl glycol, chlorphenesin, edetate disodium, glycerin, hydroxyethyl cellulose, phenoxyethanol, polysorbate 20, purified water

Principal Display Panel

CARE SCIENCE

Antibacterial

Lidocaine Wound Gel

Lidocaine Hydrochloride 2%

Benzalkonium Chloride 0.13%

Pain Relief

Antibacterial gel helps prevent infection

For treating minor cuts, scrapes and burns

Helps reduce scarring

Water-based clear hydrogel

Antibiotic free

ANTIBACTERIAL LIDOCAINE WOUND GEL

benzalkonium chloride, lidocaine hydrochloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51142-652(NDC:59898-950)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	2 g in 100 mL
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Product Characteristics

Color	white (colorless, clear translucent)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51142-652-14	1 in 1 CARTON	10/28/2019	
1		14 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:51142-652-28	1 in 1 CARTON	10/28/2019	
2		28 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/28/2019	

Labeler - ASO LLC (152793493)

Revised: 12/2022 ASO LLC