FIRST AID ANTISEPTIC- benzalkonium 0.13%, lidocaine hcl 2.5% spray
Dollar General

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.

Drug Facts

Active ingredients	**Purpose**
Benzalkonium 0.13%	First Aid Antiseptic
Lidocaine HCl 2.5%	Pain Relieving Spray

Uses

First aid to help prevent bacterial contamination or skin infection, and for temporary relief of pain and itching associated with minor: cuts, scrapes, burns, sunburn, skin irritations

Warnings

For external use only

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

When using this product

- do not use in or near the eyes
- do not apply over large area of the body or in large quantities
- do not apply over raw surfaces or blistered areas

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 right away

Directions

• Adults and children 2 years and older: clean the affected area. Apply small amount on the area 1-3 times daily; may be covered with a sterile bandage (let dry first) • Children Under 2 years: Consult a physician.

Other information

Avoid excessive heat

Inactive ingredients

Diazolidinyl Urea, Disodium EDTA, Fragrance, Methylparaben, Nonoxynol 9, Propylene Glycol, Propylparaben, Sodium Phosphate Dibasic, Water

Questions? 888-309-9030

FIRST AID ANTISEPTIC

benzalkonium 0.13%, lidocaine hcl 2.5% spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-781
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	25 mg in 1 mL

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Diazolidinyl Urea (UNII: H5RIZ3MPW4)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
Methylparaben (UNII: A2I8C7HI9T)	
NONOXYNOL-9 (UNII: 48Q180SH9T)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Propylparaben (UNII: Z8IX2SC1OH)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
Water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-781-05	148 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/29/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	01/02/2013	

Labeler - Dollar General (068331990)

Registrant - Product Quest Mfg (927768135)

Establishment

Name	Address	ID/FEI	Business Operations
Product Quest Mfg		927768135	manufacture(55910-781) , label(55910-781)

Revised: 7/2018 Dollar General