# DG HEALTH MAXIMUM STRENGTH PAIN RELIEVING CLEANSING- benzalkonium cl, lidocaine hcl spray Dollar General Corporation

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.

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#### **Drug Facts**

#### **Active ingredients**

Benzalkonium CI 0.13%

Lidocaine HCI 4%

#### **Purpose**

Benzalkonium CI - First Aid Antiseptic

Lidocaine HCI - 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 areas 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; spray a small amount on the area 1-3 times daily; may be covered with a sterile bandage (let dry first)
- children under 2 years: ask a doctor

## **Other Safety Information**

- store at room temperature
- avoid excessive heat

### **Inactive ingredients**

chlorphenesin, edetate disodium, fragrance, glycerin, nonoxynol 9, phenoxyethanol, propylene glycol, purified water

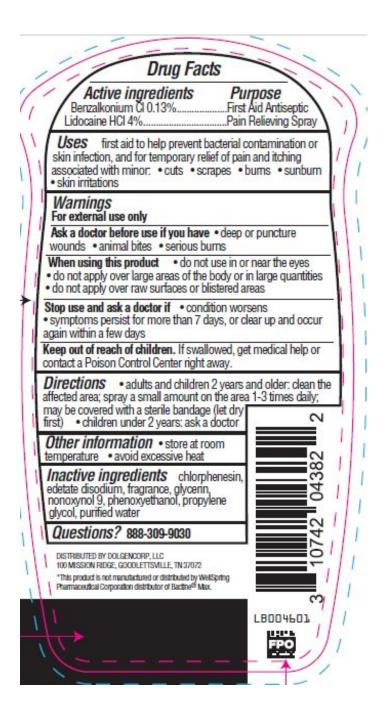
Questions

888-309-9030

Package/Label Principal Display Panel



**Principal Display Panel** 



#### DG HEALTH MAXIMUM STRENGTH PAIN RELIEVING CLEANSING

benzalkonium cl, lidocaine hcl spray

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75712-200	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL		
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	40 mg in 1 mL		

Inactive Ingredients  Ingredient Name  CHLORPHENESIN (UNII: 1670DAL4SZ)  Strength
CHLORPHENESIN (UNII: 1670DAL4SZ)
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EDETATE DISODIUM (UNII: 7FLD91C86K)
GLYCERIN (UNII: PDC6A3C0OX)
NONOXYNOL-9 (UNII: 48Q180SH9T)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
WATER (UNII: 059QF0KO0R)

l	P	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1	NDC:75712-200- 01	148 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/11/2022		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	02/11/2022		

## **Labeler -** Dollar General Corporation (006946172)

Revised: 2/2023 Dollar General Corporation