FIRST AID ANTISEPTIC PAIN RELIEVING- benzalkonium chloride, lidocaine hydrochloride cream GFA Production (Xiamen) Co., Ltd.

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First Aid Antiseptic Pain Relieving Cream

#### **Drug Facts**

#### Active ingredients

Benzalkonium chloride 0.13%

Lidocaine hydrochloride 0.5

### Purpose

First aid antiseptic

Pain relieving cream

#### Uses

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

## Warnings

#### For external use only.

#### Do not use

in the eyes
over large areas of the body
in large quantities
over raw surfaces or blistered areas
longer than 1 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 or gets worse • 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

- Clean the affected area
- Adults and children 2 years of age and older: Apply a small amount of this product to affected area not more than 3 times daily
- Children under 2 years of age: consult a doctor
- May be covered with a sterile bandage

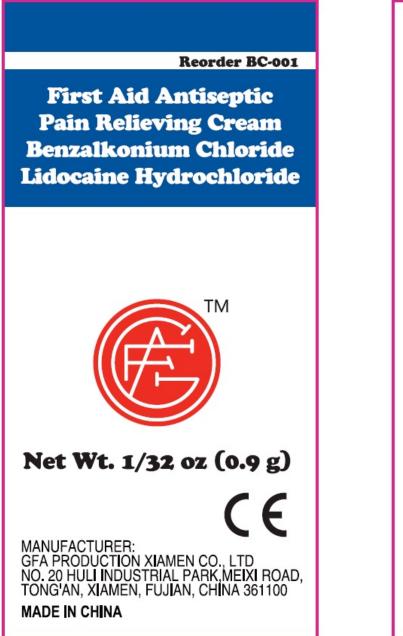
# Other information

Store at room temperature

# Inactive ingredients

glycerin monostearate, glycerol, purified water

# Package Labeling:



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#### FIRST AID ANTISEPTIC PAIN RELIEVING benzalkonium chloride, lidocaine hydrochloride cream **Product Information** HUMAN OTC DRUG NDC:50814-009 **Product Type** Item Code (Source) TOPICAL **Route of Administration Active Ingredient/Active Moiety Ingredient Name Basis of Strength** Strength BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -1.3 mg **BENZALKONIUM CHLORIDE** UNII:7N6JUD5X6Y) in 1 g LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE -LIDOCAINE HYDROCHLORIDE 5 mg UNII:98PI200987) ANHYDROUS in 1 g Inactive Ingredients **Ingredient Name** Strength **GLYCERIN** (UNII: PDC6A3C0OX) WATER (UNII: 059QF0KO0R) Packaging Marketing Start **Marketing End Package Description Item Code** # Date Date 1 NDC:50814-009- 0.9 g in 1 PACKAGE; Type 0: Not a Combination 07/01/2016 01 Product **Marketing Information Application Number or Monograph Marketing End** Marketing **Marketing Start** Category Citation Date Date OTC Monograph Drug M003 07/01/2016

Labeler - GFA Production (Xiamen) Co., Ltd. (421256261)

Revised: 1/2024

GFA Production (Xiamen) Co., Ltd.