# FIRST AID AND BURN- benzalkonium chloride and lidocaine hydrochloride cream Acme United 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 Chloride 0.13%

Lidocaine HCl 0.5%

#### **Purpose**

First Aid Antiseptic

External analgesic

#### Uses

first aid to help prevent infection and for the temporary relief of itching associated with

- minor cuts
- scrapes
- burns

#### Warnings

#### For external use only

#### Do not use

- in the eyes or apply over large areas of the body
- longer than 1 week unless directed by a doctor
- in large quantities, particularly over raw surfaces or blistered areas

Ask a doctor before use if you have deep or puncture wounds, animal bites, or serious burns

When using this product avoid contact with eyes

#### Stop use and ask a doctor if

- condition worsens or persists for more than 7 days
- clears up and occurs 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 of age and older:
- clean the affected area
- apply a small amount of this product to the area 1 to 3 times daily
- may be covered with a sterile bandage
- children under 2 years of age: consult a doctor

#### Other information

- store at room temperature
- do not use if packet is opened or torn

### **Inactive ingredients**

aloe vera, emulsifying wax, ethyl alcohol, methylparaben, mineral oil, paraffin, propylparaben, purified water, white petrolatum, white wax

Questions? 1-800-835-2263

REORDER www.Physicians CareFirstAid.com 1 800 835 2263

#### PRINCIPAL DISPLAY PANEL – 0.9 gram packet

NDC 0924-5701-01

**Physicians CARE** 

**Burn Cream** 

0.9 g (1/32 oz.)

Acme United Corporation

Fairfield, CT 06824 800-835-2263

Physicians CARE.

# Burn Cream 0.9 g (1/32 oz.)

Acme United Corporation Fairfield, CT 06824 **800-835-2263** 

# **Drug Facts**

Active ingredient Purpose
Benzalkonium First Aid
Chloride 0.13% . . . Antiseptic
Lidocaine Topical
HCl 0.5% . . . . . Analgesic

**Uses** ■ temporary relief of pain associated with minor cuts, scrapes, burns ■ helps protect against harmful bacteria

Warnings
For external use only



Warnings (continued)
Do not use ■ in eyes ■ in large quantities ■ over raw or blistered areas, or on deep puncture wounds, animal bites, serious burns ■ for more than one week unless directed by a doctor Keep out of reach of children If ingested contact a Poison Control Center right away.

Directions ■ clean affected area ■ apply small amount not more than 3 times daily ■ may be covered with a sterile bandage.

Inactive ingredients
aloe vera, emulsifying wax,
ethyl alcohol, methylparaben,
mineral oil, paraffin,
propylparaben, purified water,
white petrolatum, white wax

#### PRINCIPAL DISPLAY PANEL - 10 count box

#### **Physicians CARE**

#### **Burn Cream**

Benzalkonium chloride First aid antiseptic Lidocaine HCl External Analgesic

10 Single use Packets 0.9 g each

Reorder No. 51014

ANSI Z308.1

#### **Drug Facts**

#### Active ingredients (In each gram)

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#### FIRST AID AND BURN

benzalkonium chloride, lidocaine hydrochloride cream

#### **Product Information**

Product Type	HUMAN OTC DRUG LABEL	Item Code (Source)	NDC:0924-5701
Route of Administration	TOPICAL	DEA Schedule	

Active	Ingre	dient/ <i>E</i>	Active	Moiety

Ingredient Name	Basis of Strength	Strength
benzalkonium chloride (benzalkonium)	benzalkonium chloride	1.3 mg in 1 g
lidocaine hydrochloride (lidocaine)	lidocaine hydrochloride	5 mg in 1 g

Inactive Ingredients	
Ingredient Name	Strength
aloe vera leaf	
alcohol	
methylparaben	
mineral oil	
paraffin	
propylparaben	
water	
petrolatum	
white wax	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-5701-04	10 in 1 BOX		
1	NDC:0924-5701-01	0.9 g in 1 PACKET		
2	NDC:0924-5701-09	10 in 1 BOX		
2	NDC:0924-5701-01	0.9 g in 1 PACKET		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	0 1/27/20 12	

## Labeler - Acme United Corporation (001180207)

### **Registrant -** Safetec of America, Inc. (874965262)

Establishment				
Name	Address	ID/FEI	Business Operations	
Safetec of America, Inc.		874965262	MANUFACTURE(0924-5701)	

Revised: 2/2013 Acme United Corporation