

FIRST AID AND BURN- benzalkonium chloride, 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.

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.PhysiciansCareFirstAid.com****1 800 835 2263****Principal Display Panel - Packet Label****FIRST AID ONLY[®]****Burn Cream**

NET WT 1/32 oz (0.9g)

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Faifield, CT 06824

1.800.835.2263

www.FirstAidOnly.com

NDC 0924-5701-01



Burn Cream

NET WT 1/32 oz (0.9g)

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NDC 0924-5701-01

Drug Facts	
Active Ingredients	Purpose
Benzalkonium Chloride 0.13%	First Aid Antiseptic
Lidocaine HCl 0.5%	Topical Analgesic

Drug Facts (cont'd)

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

Warnings For external use only

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 a small amount 1 to 3 times daily ■ may be covered with a sterile bandage

Inactive Ingredient

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

Principal Display Panel - Packet Label

FIRST AID ONLY®

13-006

BURNS

First Aid/Burn Cream

12 (0.9g) Packets

Ungüento de primeros auxilios/para quemaduras

12 sobres de (0.9g)

NDC 0924-5701-10



FIRST AID AND BURN

benzalkonium chloride, lidocaine hydrochloride cream

Product Information

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

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
benzalkonium chloride (UNII: F5UM2KM3W7) (benzalkonium - UNII:7N6JUD5X6Y)	benzalkonium chloride	1.3 mg in 1 g
lidocaine hydrochloride (UNII: V13007Z41A) (lidocaine - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
aloe vera leaf (UNII: ZY81Z83H0X)	
alcohol (UNII: 3K9958V90M)	
methylparaben (UNII: A2I8C7HI9T)	
mineral oil (UNII: T5L8T28FGP)	
paraffin (UNII: I9O0E3H2ZE)	
propylparaben (UNII: Z8IX2SC10H)	
water (UNII: 059QF0KOOR)	
petrolatum (UNII: 4T6H12BN9U)	
white wax (UNII: 7G1J5DA97F)	

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; Type 0: Not a Combination Product		
2	NDC:0924-5701-09	10 in 1 BOX		
2	NDC:0924-5701-01	0.9 g in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:0924-5701-10	12 in 1 BOX		
3	NDC:0924-5701-01	0.9 g in 1 PACKET; Type 0: Not a Combination Product		
4	NDC:0924-5701-11	25 in 1 BOX		
4	NDC:0924-5701-01	0.9 g in 1 PACKET; Type 0: Not a Combination Product		
5	NDC:0924-5701-12	144 in 1 BOX		
5	NDC:0924-5701-01	0.9 g in 1 PACKET; Type 0: Not a Combination Product		
6	NDC:0924-5701-13	60 in 1 BOX		
6	NDC:0924-5701-01	0.9 g in 1 PACKET; 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	01/27/2012	

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: 3/2016

Acme United Corporation