FIRST AID ONLY FIRST AID-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.

First Aid Only First Aid-Burn Cream

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 pain and itching associated with minor

- cuts
- scrapes
- burns.

Warnings

For external use only

Do not use

- in the eyes
- over large areas of the body
- in large quantities, particularly over raw surfaces or blistered areas
- if you are allergic to any of the ingredients
- on deep puncture wounds, animal bites, or serious burns
- more than one week unless directed by a doctor

Keep out of reach of children.

If swallowed contact a Poison Control Center right away.

Directions

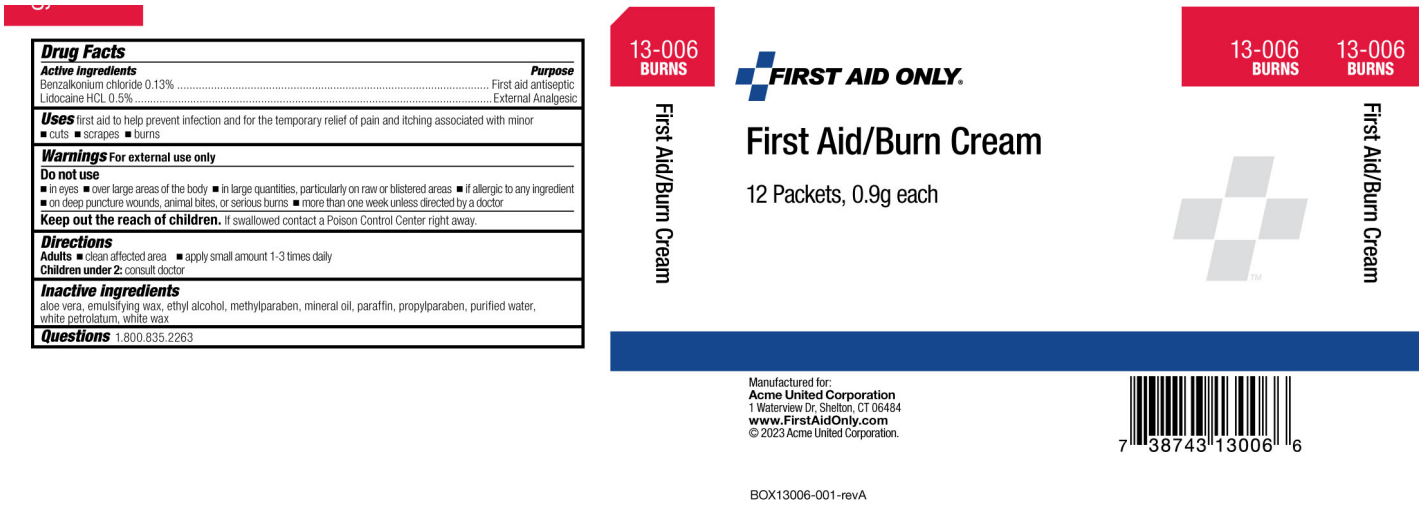
Adults

- clean affected area
- apply small amount 1-3 times daily
- **children under 2**: consult doctor

Inactive ingredients

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

Box Label



FIRST AID ONLY FIRST AID-BURN

benzalkonium chloride, lidocaine hydrochloride cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0924-5702(NDC:61010-5701)
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: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
PETROLATUM (UNII: 4T6H12BN9U)	
WHITE WAX (UNII: 7G1J5DA97F)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-5702-00	0.9 g in 1 PACKET; Type 0: Not a Combination Product	01/30/2023	
2	NDC:0924-5702-01	6 in 1 BOX	08/14/2023	
2		0.9 g in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:0924-5702-02	10 in 1 BOX	08/14/2023	
3		0.9 g in 1 PACKET; Type 0: Not a Combination Product		
4	NDC:0924-5702-03	12 in 1 BOX	08/14/2023	
4		0.9 g in 1 PACKET; Type 0: Not a Combination Product		
5	NDC:0924-5702-04	20 in 1 BOX	08/14/2023	
5		0.9 g in 1 PACKET; Type 0: Not a Combination Product		
6	NDC:0924-5702-05	25 in 1 BOX	08/14/2023	
6		0.9 g in 1 PACKET; Type 0: Not a Combination Product		
7	NDC:0924-5702-06	60 in 1 BOX	08/14/2023	
7		0.9 g in 1 PACKET; Type 0: Not a Combination Product		
8	NDC:0924-5702-07	144 in 1 BOX	08/14/2023	
8		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	01/30/2023	

Labeler - Acme United Corporation (001180207)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	repack(0924-5702) , relabel(0924-5702)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	relabel(0924-5702) , repack(0924-5702)