FIRST AID AND BURN- benzalkonium chloride, lidocaine cream HART Health

FIRST AID & BURN CREAM

Active Ingredients (in each gram):

Benzalkonium Chloride 0.13%

Lidocaine HCI 0.5%

Purpose:

Topical Antiseptic

Topical Anesthetic

Uses: Temporary relief of pain and to help prevent infection in

- minor cuts
- scrapes
- burns

Warnings: For external use only. See a doctor immediately for serious burns

Do not use:

- in or near the eyes
- over large portions of the body
- in large quantities particularly over raw surfaces or blistered areas
- longer than 7 days unless directed by a doctor
- more than 1 to 3 times daily

Ask a doctor before use if you have

- serious burns
- animal bites
- deep or puncture wounds

Stop use and ask a doctor if

- condition lasts for more than 7 days
- condition worsens
- symptoms 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. 1-800-222-1222

Directions: do not use more than directed

Adults and children 2 years of age and over:

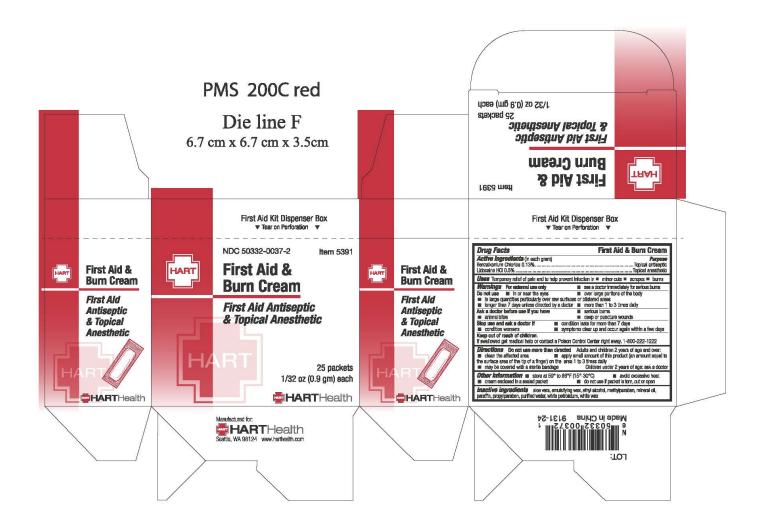
- clean the affected area
- apply small amount of this product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily
- may be covered with a sterile bandage

Children under 2 years of age: ask a doctor

Inactive Ingredients: Aloe Vera, Cetyl Alcohol, Diazolidinyl Urea, Edetate Disodium, Glycerin, Clyceryl Monostearate, Methylparaben, Mineral Oil, Polyethylene Glycol, Propylene Glycol, Propylparaben, Purified Water, Stearic Acid, Trolamine







FIRST AID AND BURN

benzalkonium chloride, lidocaine cream

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:50332-0037

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y) BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - CHLORIDE in 100 g LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987) LIDOCAINE (UNII: 98PI200987)

Inactive Ingredients				
Ingredient Name	Strength			
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
ALCOHOL (UNII: 3K9958V90M)				
METHYLPARABEN (UNII: A2I8C7HI9T)				
MINERAL OIL (UNII: T5L8T28FGP)				

PARAFFIN (UNII: 1900E3H2ZE)	
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:50332- 0037-1	6 in 1 BOX, UNIT-DOSE	07/14/2000			
1		.9 g in 1 PACKET; Type 0: Not a Combination Product				
2	NDC:50332- 0037-6	10 in 1 BOX, UNIT-DOSE	07/14/2000			
2		.9 g in 1 PACKET; Type 0: Not a Combination Product				
3	NDC:50332- 0037-2	25 in 1 BOX, UNIT-DOSE	07/14/2000			
3		.9 g in 1 PACKET; Type 0: Not a Combination Product				
4	NDC:50332- 0037-4	144 in 1 BOX, UNIT-DOSE	07/14/2000			
4		.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 Drug	M	07/14/2000			

Labeler - HART Health (069560969)

Revised: 2/2024 HART Health