MEDI-FIRST PLUS BURN CREAM- benzalkonium chloride, lidocaine hcl cream
Unifirst First Aid Corporation

Medi-First Plus Burn Cream

Drug Facts

Active ingredients

Benzalkonium Chloride 0.13%

Lidocaine HCl 0.5%

Purpose

First aid antiseptic

Topical analgesic

Uses

- for the temporary relief of pain associated with minor burns
- helps protect against harmful bacteria

Warnings

For external use only.

Do not use

- in the eyes
- in large quantities
- over raw or blistered areasover large areas, or on deep puncture wounds, animal bites, or serious burns
- for more than 1 week unless directed by a doctor

Stop use and ask a doctor if

- the condition gets worsens or if symptoms persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

Ifingested, get medical help or contact a Poison Control Center right away.

Directions

- **Adults and children 12 years and over:**

clean the affected area, apply a small amount not more than 3 to 4 times daily

- **Children under 12 years:**

do not use, consult a doctor

Other information

- store at room temperature (do not freeze)
- tamper evident sealed packets
- do not use any opened or torn packets

Inactive ingredients

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

Questions or comments? 800-634-7680

Medi-First Plus Burn Cream Label

Medi First®

Plus

Burn Cream

with Lidocaine

Benzalkonium Chloride 0.13% / Lidocaine HCl 0.5%

Topical Antiseptic / Topical Analgesic

25 Units of 0.9g Packets

Item #93973

MEDI-FIRST PLUS BURN CREAM

benzalkonium chloride, lidocaine hcl cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47682-940
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	5 mg in 1 g
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 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)	
PETROLATUM (UNII: 4T6H12BN9U)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
WHITE WAX (UNII: 7G1J5DA97F)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-940-73	25 in 1 BOX	08/29/2017	
1		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 Drug	M003	08/29/2017	

Labeler - Unifirst First Aid Corporation (832947092)

Revised: 1/2024

Unifirst First Aid Corporation