

**BURN TREATMENT- benzalkonium chloride, lidocaine hydrochloride cream  
CMC Group Inc.**

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**Burn Treatment**

***Drug Facts***

***Active ingredients***

Benzalkonium chloride 0.13%

Lidocaine hydrochloride 0.5%

**Purpose**

First aid antiseptic

Pain relieving cream

**Uses**

- First aid to help prevent infection in minor cuts, scrapes, and burns.
- For the temporary relief of pain and itching associated with minor burns , minor cuts, and scrapes

***Warnings***

For external use only.

***Do not use***

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

**Ask a doctor before use if you have**

- deep or puncture wounds • animal bites • serious burns.

**Stop use and ask a doctor if**

- the condition persists or gets worse • symptoms persist for more than 7 days or 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.

***Directions***

- Clean the affected area

- Adults and children 2 years of age and older: Apply a small amount of this product to affected area not more than 3 times daily
- Children under 2 years of age: consult a doctor
- May be covered with a sterile bandage

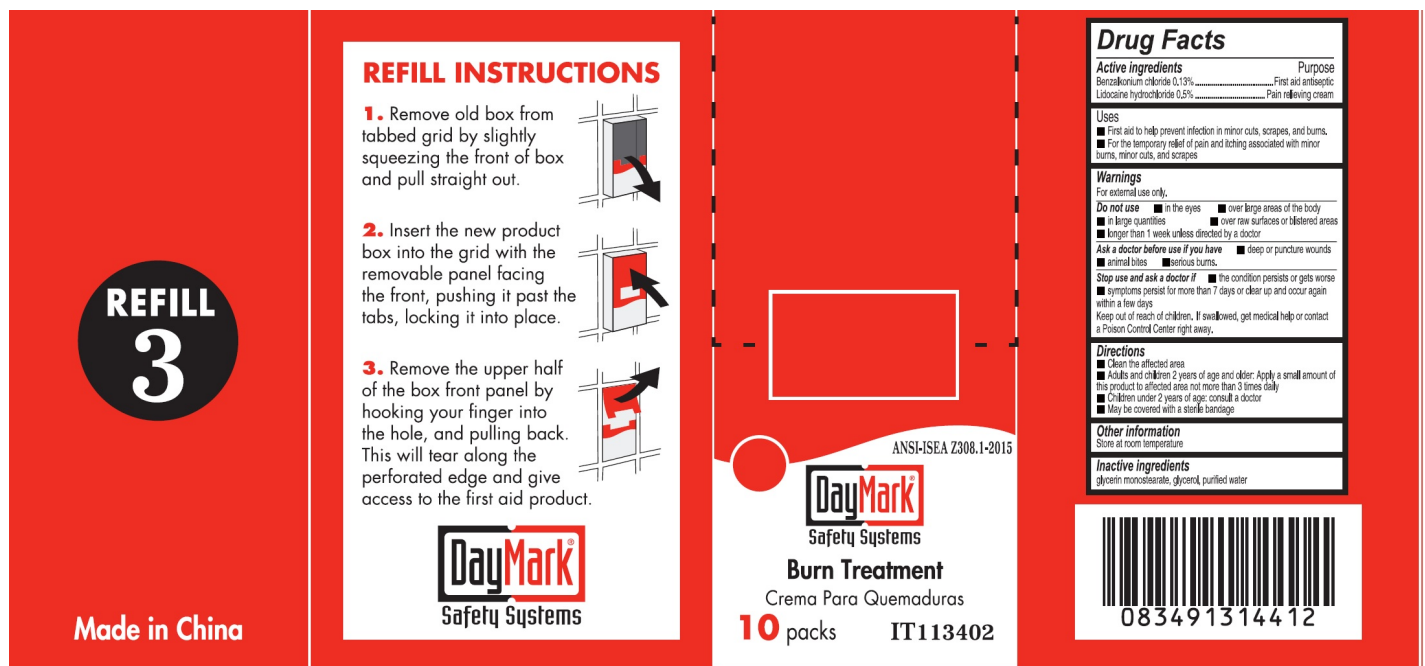
### Other information

Store at room temperature

### Inactive ingredients

glycerin monostearate, glycerol, purified water

### Package Labeling:



## BURN TREATMENT

benzalkonium chloride, lidocaine hydrochloride cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:49687-0014
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII: 98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>WATER</b> (UNII: 059QF0KO0R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49687-0014-0	10 in 1 KIT	08/06/2016	
1		0.9 g in 1 POUCH; 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	08/06/2016	

**Labeler** - CMC Group Inc. (117201448)

Revised: 1/2025

CMC Group Inc.