

BURN CREAM- benzalkonium chloride, lidocaine hydrochloride cream
Total Resources International

Drug Facts

Active Ingredients

Benzalkonium Chloride 0.13% w/w

Lidocaine HCl 0.5% w/w

Purposes

First aid antiseptic

Topical analgesic

Uses

- temporary relief of pain associated with minor 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 small amount not more than 3 times daily
- children under 2: consult a doctor

Inactive Ingredients

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

Principal Display Panel - 0.03 oz Pouch Label

BE SMART

GET PREPARED®

BURN CREAM

0.03 oz [0.90 g]

Single Use

Made in USA

Manufactured for Total Resources Intl. Inc.

Walnut, California 91789

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NDC 55550-902-21



BURN CREAM

benzalkonium chloride, lidocaine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55550-902
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:55550-902-21	0.9 g in 1 POUCH; Type 0: Not a Combination Product	01/14/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/14/2020	

Labeler - Total Resources International (790160535)