BE SMART GET PREPARED BURN CREAM- benzalkonium chloride, lidocaine hydrochloride anhydrous cream
Total Resources International, Inc

Drug Facts

Active Ingredients

Benzalkonium Chloride 0.13% w/w

Lidocaine HCl 0.5% w/w

Purpose

First Aid Antiseptic

Topical Analgesic

Uses

- Temporary relief of pain associated with minor cuts, scrapes, and 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, or serious burns
- for more than one week unless directed by a doctor

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center directly.

Directions

- clean affected area
- apply small amount not more than 3 times daily
- may be covered with a sterile bandage

Other Information

- store at room temperature
- tamper evident sealed packets

- do not use any opened or torn packets

Inactive Ingredients

aloe barbadensis leaf juice, cetearyl alcohol, disodium edta, ethylhexylglycerin, glycerin, glyceryl stearate/peg-100 stearate, maltodextrin, mineral oil, phenoxyethanol, propylene glycol, stearic acid, sodium hydroxide, water

Principal Display Panel – 0.90mL Pouch Label

BE SMART

GET PREPARED®

BURN CREAM

0.03 oz [0.90 g]

Single Use

Made in China

Manufactured for Total Resources Intl. Inc.

Walnut, California 91789

www.besmartgetprepared.com • *00-BUR-0011COM Rev. 00*

NDC 55550-521-01

BE SMART GET PREPARED BURN CREAM

benzalkonium chloride, lidocaine hydrochloride anhydrous cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55550-521
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

LIDOCAINE HYDROCHLORIDE ANHYDROUS (UNII: EC2CNF7XFP) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERYL STEARATE/PEG-100 STEARATE (UNII: RD25J5V947)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
MINERAL OIL (UNII: T5L8T28FGP)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55550-521-01	0.9 g in 1 POUCH; Type 0: Not a Combination Product	08/16/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	08/16/2024	

Labeler - Total Resources International, Inc (790160535)

Revised: 8/2024 Total Resources International, Inc