FIRST AID BURN CREAM- first aid burn cream cream
Front Line Safety

Active Ingredient

Lidocaine Hydrochloride 0.5%

Purpose

Topical Analgestic

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Topical Antiseptic

Use(s)

• For the temporary relief of pain and itching associated with:

• Sunburn • Minor burns • Insect bites • Minor skin irritations • Cuts • Scrapes

Warnings

For External Use Only

Do not use

• In the eyes • Over large areas of the body or on deep puncture wounds, animal bites, or serious burns

• In large quantities, particularly over raw surfaces or blistered areas

Stop use and aks a doctor if

• The condition get worse • Condition clears up and recurs within a few days • Condition persists for more than 7 days

If pregnant or breast feeding

Ask a health care prfessional before use.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222)

right away.

Directions

Adults and children 2 years and over:

- Clean the affected area
- Apply a small amount of this product on the area 3 to 4 times daily
- May be covered with a sterile bandage

Children under 2 years: • Consult a doctor

Other Information

- Store in a cool, dry area 15º-25ºC (59º-77ºF)
- Tamper evident sealed packets
- Do not use any opened or torn packets

Inactive Ingredients

Butylated Hydroxytoluene, Cetomacrogol 1000, Cetostearyl Alcohol, Dimethicone, Glycerine, Glyceryl Monostearate, Isopropyl Myristate, Methylcellulose, Purified Water, Sodium EDTA, Sodium Methylparaben, Sodium Propylparaben

Questions?

1-888-900-2920 Monday - Friday 8AM-4PM PST

Label

FL-2044

FIRST AID BURN CREAM

first aid burn cream cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58228-6244
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	50 mg in 10000 mg
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	13 mg in 10000 mg

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLPARABEN SODIUM (UNII: 625NNB0G9N)	
CETETH-20 (UNII: I835H2IHHX)	
METHYLPARABEN SODIUM (UNII: CR6K9C2NHK)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	

CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
WATER (UNII: 059QF0KO0R)	
EDETATE SODIUM (UNII: MP1J8420LU)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
METHYLCELLULOSE (4000 MPA.S) (UNII: MRJ667KA5E)	
DIMETHICONE 350 (UNII: 2Y53S6ATLU)	

Product Characteristics

Color		**Score**	
Shape	FREEFORM	**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58228-6244-1	1800 in 1 CASE	01/26/2024	
1		25 in 1 BOX		
1		900 mg 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	M017	01/26/2024	

Labeler - Front Line Safety (061263699)

Revised: 11/2024 Front Line Safety