BURN EASE- burn ease gel
Front Line Safety

FL-8610

Active Ingredient

Lidocaine Hydrochloride USP 2%

Purpose

Analgesic

Use(s)

For the temporary relief of pain associated with • Minor burns • Sunburn

Warnings

For External Use Only

Do not use

• On wounds or damaged skin

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

When using this product

• Avoid contact with the eyes

• Do not bandage tightly

Stop use and ask a doctor if

• Condition worsens

• Symptoms persists for more than 7 days

• Symptoms 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 (1-800-222-1222) right away

Directions

• Adults and children 2 year of age and older: Apply to affected area not more than 3 to 4 times daily

• Children under 2 years of age: Consult a doctor

Other Infomation

• Store at controlled room temperature 20º-25ºC (68º-77ºF)

• Tamper Evident. Do not use if seal is damaged.

Inactive Ingredients

Acrylates/C10-30 Alkyl Acrylate Crosscopolymer, Carbomer, Glycerin, Imidazolidinyl Urea, Methylparaben, Propylene Glycol, Propylparaben, Purified Water, Tea Tree Oil, Trolamine

Questions?

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

Label

BURN EASE

burn ease gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58228-6210
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 809Y72KV36)	
IMIDUREA (UNII: M629807ATL)	
TEA TREE OIL (UNII: VIF565UC2G)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
METHYLPARABEN (UNII: A2I8C7HI9T)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58228-6210-1	900 in 1 CASE	01/08/2024	
1		25 in 1 BOX		
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	M017	01/08/2024	

Labeler - Front Line Safety (061263699)

Revised: 11/2024

Front Line Safety