

BURN EASE 3.5G- burn ease 3.5g gel
Front Line Safety

FL-3832

Active Ingredient

Lidocaine Hydrochloride 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
- 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 persist 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 years of age and older:** Apply to affected area not more than 3 to 4 times daily

- **Children under 2 year of age:** Consult a doctor

Other information

- Store at room temperature

Inactive Ingredients

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

Questions?

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

Label



FL-3832

BURN EASE 3.5G			
burn ease 3.5g gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58228-6232
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: PDC6A3C00X)				
METHYLPARABEN (UNII: A2I8C7HI9T)				
WATER (UNII: 059QF0KO0R)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 809Y72KV36)				
PROPYLPARABEN (UNII: Z8IX2SC1OH)				
IMIDUREA (UNII: M629807ATL)				
TEA TREE OIL (UNII: VIF565UC2G)				
TROLAMINE (UNII: 9O3K93S3TK)				
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)				
Product Characteristics				
Color		Score		
Shape	FREEFORM	Size		
Flavor		Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58228-6232-1	600 in 1 CASE	01/26/2024	
1		6 in 1 BOX		
1		3.5 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	01/26/2024		

Labeler - Front Line Safety (061263699)