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

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**FL-3831**

### *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

• 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 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, Imidazolidinly Urea, Methylparaben, Propylene Glycol, Propylparaben, Purified Water, Tea Tree Leaf Oil, Triethanolamine

## *Questions?*

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

## *Label*


FL-3831

| **BURN EASE 3.5G** |
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| burn ease 3.5g gel |

## Product Information

| | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:58228-6231 |
| **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 |
|---|---|
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **METHYLPARABEN** (UNII: A2I8C7HI9T) | |
| **IMIDUREA** (UNII: M629807ATL) | |
| **CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED)** (UNII: 809Y72KV36) | |
| **PROPYLPARABEN** (UNII: Z8IX2SC1OH) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **TROLAMINE** (UNII: 9O3K93S3TK) | |
| **CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE** (UNII: 0A5MM307FC) | |
| **TEA TREE OIL** (UNII: VIF565UC2G) | |

## Product Characteristics

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

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:58228-6231-1 | 480 in 1 CASE | 01/26/2024 | |
| **1** | | 20 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)

Revised: 12/2025

Front Line Safety