WALGREENS SUNBURN RELIEF- lidocaine gel WALGREEN COMPANY

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Walgreens Sunburn Relief Gel

Active ingredient

Lidocaine 0.8% (as Lidocaine HCI)

Purpose

External Analgesic

Uses

For the temporary relief of pain and itching associated with sunburn, minor burns, minor cuts, scrapes, insect bites, and minor skin irritations

Warnings

For External Use Only

Do not use

in large quantities, particularly over raw surfaces or blistered areas

When using this product

avoid contact with eyes. If contact occurs, rinse with water to remove.

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, seek medical help or contact a Poison Control Center immediately.

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 years of age: Ask a doctor.

Inactive ingredients

water, aloe barbadensis leaf juice, alcohol denat., propylene glycol, laureth-23, glycerin, polysorbate 20, allantoin, carbomer, triethanolamine, menthyl lactate, disodium EDTA, menthol, aleurites moluccanus seed extract, carica papaya (papaya) fruit extract, colocasia antiquorum root extract, mangifera indica (mango) fruit extract, passiflora incarnata flower extract, plumeria acutifolia flower extract, psidium guajava fruit extract, tocopheryl acetate, tocopherol, phenoxyethanol, benzyl alcohol, fragrance, blue 1.



Walgreens Sunburn Relief Gel with lidocaine HCI relieves the pain of sunburns, minor skin irritations and abrasions. This soothing gel nourishes and helps restore moisture to dry and damaged skin.

Drug Facts

Active ingredient Lidocaine 0.8% (as Lidocaine HCI).

Purpose External Analgesic

Uses For the temporary relief of pain and itching associated with sunburn, minor burns, minor cuts, scrapes, insect bites, and minor skin irritations.

For External Use Only

Do not use in large quantities, particularly over raw surfaces or blistered areas

When using this product avoid contact with eyes, If contact occurs, rinse with water

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, seek medical help or contact a Poison Control Center immediately,

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 years of age: Ask a doctor.

Inactive ingredients: water, also barbadensis leaf juice, alcohol denat, propylene glycol, laureth-23, glycerin, polysorbate 20, allantoin, carbomer, triethanolsmine, menthyl lactate, disodium EDTA, menthol, aleurites moluccanus seed extract, carica papaya (papaya) fruit extract, colocasia antiquorum root extract, mangifera indica (mango) fruit extract, passiflora incarnata flower extract, plumeria acutifolia flower extract, psidium guajava fruit extract, tocopheryl acetate, tocopherol, phenoxyethanol, benzyl alcohol, fragrance, blue 1.

Call toll free 1-800-925-4733



Health expertise you rely on, quality you trust.

[†]Walgreens Pharmacist Survey

DISTRIBUTED BY: WALGREEN CO. 200 WILMOT RD, DEERFIELD, IL 60015 100% SATISFACTION GUARANTEED walgreens.com @2020 Walgreen Co. ITEM 991466 W10265-0620-F



ORG0717-F REV-0620

WALGREENS SUNBURN RELIEF

lidocaine gel

Product Information

Product Type HUMAN OTC DRUG

Item Code (Source)

NDC:0363-0985

Route of Administration

TOPICAL

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|-------------|
| LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987) | LIDOCAINE | 8 mg in 1 g |

| Inactive Ingredients | |
|---|----------|
| Ingredient Name | Strength |
| MANGO (UNII: 162913NR86) | |
| ALCOHOL (UNII: 3K9958V90M) | |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | |
| LAURETH-23 (UNII: N72LMW566G) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| POLYSORBATE 20 (UNII: 7T1F30V5YH) | |
| ALLANTOIN (UNII: 344S277G0Z) | |
| CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO) | |
| TROLAMINE (UNII: 903K93S3TK) | |
| MENTHYL LACTATE, (-)- (UNII: 2BF9E65L7I) | |
| EDETATE DISODIUM (UNII: 7FLD91C86K) | |
| LEVOMENTHOL (UNII: BZ1R15MTK7) | |
| KUKUI NUT OIL (UNII: TP11QR7B8R) | |
| PAPAYA (UNII: KU94FIY6JB) | |
| COLOCASIA ESCULENTA ROOT (UNII: H7B71Q0G0D) | |
| PASSIFLORA INCARNATA FLOWER (UNII: K8F3G29S6Z) | |
| GUAVA (UNII: 74070D6VG0) | |
| .ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0) | |
| .ALPHATOCOPHEROL, DL- (UNII: 7QWA1RIO01) | |
| PHENOXYETHANOL (UNII: HIE492ZZ3T) | |
| BENZYL ALCOHOL (UNII: LKG8494WBH) | |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | |
| ALOE VERA LEAF (UNII: ZY81Z83H0X) | |
| WATER (UNII: 059QF0KO0R) | |
| PLUMERIA ALBA FLOWER OIL (UNII: T69Z2432CU) | |

| Packaging | | | | | |
|-----------|----------------------|--|-------------------------|-----------------------|--|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date | |
| | NDC:0363-0985- 68 | 453 g in 1 BOTTLE; Type 0: Not a Combination Product | 11/08/2017 | | |

| Marketing Information | | | | |
|-------------------------|---|-------------------------|-----------------------|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| OTC monograph not final | part348 | 10/03/2017 | | |
| | | | | |

Revised: 1/2022 WALGREEN COMPANY