

BURN N BITE- lidocaine, menthol gel

Q.A. Laboratories

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.

Burn N Bite

Drug Facts

Active Ingredient

Lidocaine 2%

Menthol 1%

Purpose

Ant-itch, Pain Relief

Uses

for the temporary relief of pain and discomfort caused by • minor burns • sunburn • scrapes • insect bites • minor skin irritations • rashes due to poison ivy, oak and sumac.

Warnings

For external use only.

When using this product

- Avoid contact with eyes

Do not use

- Under bandages or compresses

Stop use and ask a doctor if condition worsens, or if symptoms persist for more than 7 days or 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 right away.

Directions

Adults and children 2 years of age and older: Apply to the affected area not more than two times daily.

Children under 2 years of age: Do not use, consult a doctor

Inactive Ingredients

Deionized water, propylene glycol, aloe, isopropyl alcohol, carbomer, triethanolamine, phenoxyethanol

Questions or comments:

858-782-6627, 816-421-8081

Stanmar Laboratories

Division of Q.A. Laboratories

404 Admiral Boulevard Kansas City, MO 64106

Ideazbrands.com

Made in the USA

Packaging

NDC 0520999005

DR. GREENFIELD'S
SOOTHE IT RIGHT WITH
Burn N Bite
First Aid Antiseptic Ointment

For the temporary relief of pain and discomfort caused by, minor burns, sunburn, scrapes, insect bites, minor skin irritations, rashes due to poison ivy, oak and sumac

Fast Acting - Non-Greasy
Aloe Enriched

NET WT. 5 OZ (141g)

Drug Facts	
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Lidocaine 2%.....	Anti-itch, Pain Relief
Menthol 1%.....	Anti-itch, Pain Relief

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- sunburn • scrapes • insect bites
- minor skin irritations • rashes due to poison ivy, oak and sumac

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NDC 0520999005

DR. GREENFIELD'S

SOOTHE IT RIGHT WITH

Burn N Bite

lidocaine and menthol gel

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Alone Enriched

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BURN N BITE

lidocaine, menthol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52099-9005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	2 g in 100 g
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE (UNII: V5VD430YW9)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 903K93S3TK)	
PHENOXYETHANOL (UNII: H1E492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52099-9005-1	141 g in 1 TUBE; Type 0: Not a Combination Product	03/30/1994	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	03/30/1994	

Labeler - Q.A. Laboratories (065361149)

Establishment

Name	Address	ID/FEI	Business Operations
Q.A. Laboratories		065361149	manufacture(52099-9005)

Revised: 1/2022

Q.A. Laboratories