MEDICAINE STING AND BITE- benzocaine swab Dynarex Corporation

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

1407 Medicaine Insect Bite (Ampule) 67777-405-01

active ingredient each swab has benzocaine usp, I menthol

uses for temporary relief of pain and itching associated with insect bites and stings

keep out of reach of children. not for use with children less than 2 years old without medical advice. if swallowed, get medical help immediately or contact a poison control center right away

revers cardboard sleeve then crush at dot between thumb and forefinger. once solution has saturated tip, apply topically to the sting or bit. may be used on affected area(s) up to 4 times per day

store at room temperature away from light

purified water usp

for temporary relief of pain and itching

for external use only

medicaine sting and bit ampules latex free



MEDICAINE STING AND BITE

benzocaine swab

| Product Information | | | | |
|-------------------------|----------------|--------------------|---------------|--|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:67777-405 | |
| Route of Administration | TOPICAL | | | |

| Active Ingredient/Active Moiety | | | |
|--|--------------------------|------------------|--|
| Ingredient Name | Basis of Strength | Strength | |
| BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5) | BENZOCAINE | 0.12 g in 0.6 mL | |

| Inactive Ingredients | |
|----------------------------|----------|
| Ingredient Name | Strength |
| MENTHOL (UNII: L7T10EIP3A) | |

| Packaging | | | | |
|----------------|---|-------------------------|-----------------------|--|
| # Item Code | Package Description | Marketing Start Date | Marketing End Date | |
| NDC:67777-405- | 10 in 1 BOX | 02/14/1976 | | |
| NDC:67777-405- | 6 mL in 1 AMPULE; Type 0: Not a Combination Product | | | |

| Marketing Information | | | | |
|-----------------------|---|-------------------------|-----------------------|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| unapproved drug other | | 02/14/1976 | | |
| | | | | |

Labeler - Dynarex Corporation (008124539)

Registrant - Dynarex Corporation (008124529)

Revised: 11/2022 Dynarex Corporation