

STING RELIEF PAD- benzocaine, isopropyl alcohol swab
Acme United Corporation

First Aid Only Sting Relief Pad

Drug Facts

Active Ingredients

Benzocaine, 6%

Isopropyl alcohol 60% w/v

Purpose

Topical Analgesic

Antiseptic

Use

For the temporary relief of pain and itching associated with minor burns, scrapes and insect bites. First aid to help prevent infection in minor cuts, scrapes and burns.

Warnings

For external use only

Flammable, keep away from fire or flame

Do Not use

- in eyes, if contact occurs flush with water
- over large areas of the body

Keep out of the reach of children.

If swallowed get medical help or contact a poison control center right away.

Directions

- for adults and children 2 years of age or older, apply to affected area not more than 3 to 4 times daily
- Children under 2 years: consult physician

Other Information

Store at room temperature

Inactive Ingredients

purified water

Package Labeling:



STING RELIEF PAD

benzocaine, isopropyl alcohol swab

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|------------------------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:0924-5202(NDC:59050-059) |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-------------------|---------------|
| BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5) | BENZOCAINE | 60 mg in 1 g |
| ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302) | ISOPROPYL ALCOHOL | 600 mg in 1 g |

Inactive Ingredients

| Ingredient Name | Strength |
|---------------------------------|----------|
| WATER (UNII: 059QF0KO0R) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:0924-5202-01 | 0.34 g in 1 POUCH; Type 0: Not a Combination Product | 05/09/2018 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M017 | 05/09/2018 | |

Labeler - Acme United Corporation (001180207)

Registrant - Acme United Corporation (001180207)

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|--|
| Acme United Corporation | | 045924339 | relabel(0924-5202) , repack(0924-5202) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|--|
| Acme United Corporation | | 080119599 | repack(0924-5202) , relabel(0924-5202) |

Revised: 11/2024

Acme United Corporation