FIRST AID ONLY STING RELIEF PAD- benzocaine swab
Acme United Corporation

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.

First Aid Only Sting Relief Pad

Active Ingredients

Benzocaine, 6% w/v

Isopropyl Alcohol 60% w/v

Purpose

Topical Anesthetic

Antiseptic

Use

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

Directions

▪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 a physician

Warnings

For external use only

Flammable, keep away from fire or flames

Do not use

- In the eyes. If contact occurs, rinse thoroughly with water
- over large areas of the body

Stop use

If irritation, redness or other symptoms develop. Consult a doctor if the condition persists or gets worse.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Inactive Ingredients

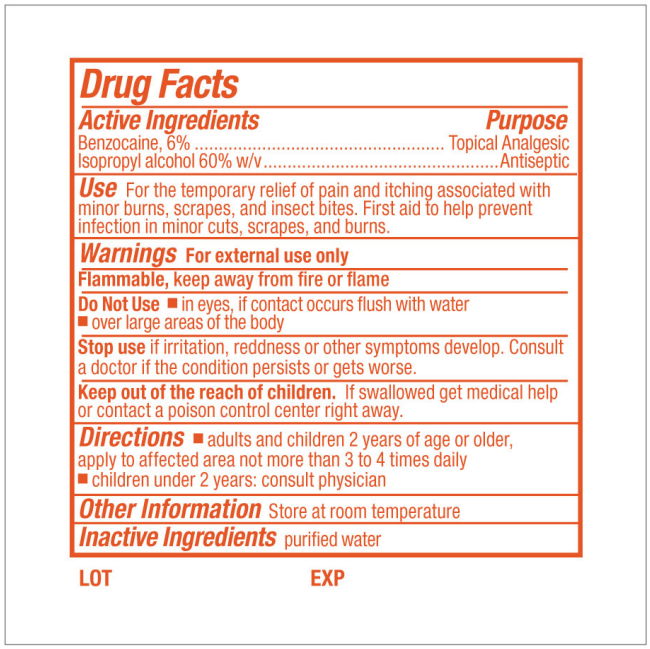
Purified Water

Other Information

Store at room temperature

Principal Display Panel - 0.4 mL Pouch Label

0.4 mL Pouch Label



FIRST AID ONLY STING RELIEF PAD

benzocaine swab

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0924-5203(NDC:65517-0005)
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 mL
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.6 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-5203-01	0.4 mL in 1 POUCH; Type 0: Not a Combination Product	04/08/2019	
2	NDC:0924-5203-02	10 in 1 BOX	04/08/2019	
2		0.4 mL in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:0924-5203-03	25 in 1 BOX	04/08/2019	
3	NDC:0924-5203-01	0.4 mL in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:0924-5203-04	100 in 1 BOX	04/08/2019	

4	NDC:0924-5203-01	0.4 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	04/08/2019	

Labeler - Acme United Corporation (001180207)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	relabel(0924-5203) , repack(0924-5203)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	relabel(0924-5203) , repack(0924-5203)

Revised: 4/2019 Acme United Corporation