FIRST AID ONLY STING RELIEF PAD- benzocaine, isopropyl alcohol liquid
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

Drug Facts

Active ingredients

Benzocaine 6.0%

Isopropyl Alcohol 60% w/v

Purpose

Topical Anesthetic

Antiseptic

For the temporary relief of pain and itching associated with minor 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**

Consult a doctor ■if condition worsens ■if symptoms last for more than seven days or clear up and occur again ■for deep puncture wounds

Keep out of reach of children.

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

Directions

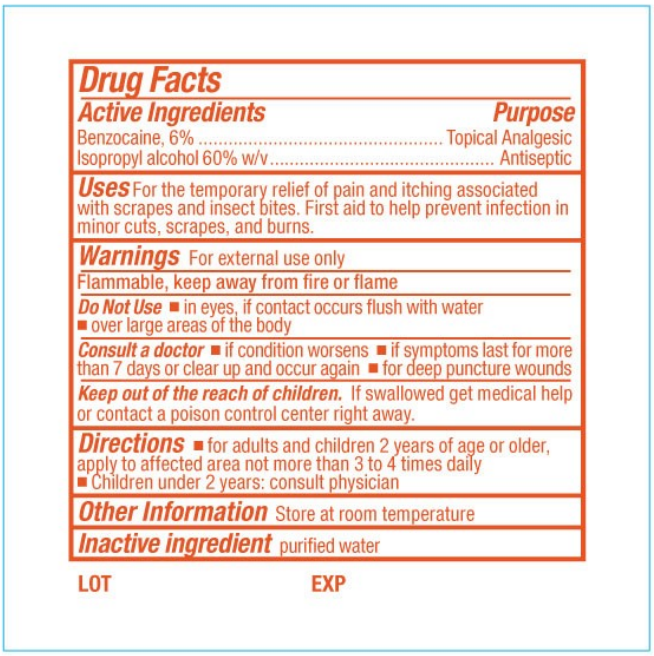
■ For adults and children 2 years of age and older, apply to affected area not more than 3 to 4 times daily

■Children under 2 years: consult physician

Inactive ingredient

Purified Water

Other information store at room temperature



pouch label

FIRST AID ONLY STING RELIEF PAD

benzocaine, isopropyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0924-5204(NDC:59050-414)
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	600 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-5204-05	100 in 1 BOX	02/25/2022	
1		0.4 mL in 1 POUCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		
2	NDC:0924-5204-04	50 in 1 BOX	02/25/2022	
2		0.4 mL in 1 POUCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		
3	NDC:0924-5204-03	25 in 1 BOX	02/25/2022	
3		0.4 mL in 1 POUCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		
4	NDC:0924-5204-02	10 in 1 BOX	02/25/2022	
4		0.4 mL in 1 POUCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		
5	NDC:0924-5204-01	0.4 mL in 1 POUCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)	02/25/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/25/2022	

Labeler - Acme United Corporation (001180207)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	relabel(0924-5204) , repack(0924-5204)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	relabel(0924-5204) , repack(0924-5204)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		117825595	manufacture(0924-5204)