FIRST AID ONLY STING RELIEF- benzocaine swab Acme United 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.

First Aid Only Sting Relief Swabs

Benzocaine 20 percent

Menthol 1 percent

Topical Analgesic

Topical Analgesic

Uses

For temporary relief of the pain and itching of

- •insect bites and stings
- minor scrapes and burns

Warnings

For external use only.

Do not use

•in the eyes •if you are allergic to any of the ingredients

Stop use and ask a doctor if •conditions worsen •symptoms clear up and then recur •the condition persists for more than 7 days

Keep out of the reach of children

If swallowed, contact Poison Control 800.222.1222, or a doctor immediately.

Directions If stinger is present, remove carefully if possible.

Adults Reverse cardboard sleeve so swab is visible. With cardboard sleeve on ampile, crush at dot on sleeve.

There will be a "pop" and the swab will become saturated. Apply saturated applicator to the affected area no more than

3 to 4 times daily.

Children under 2 Consult a doctor.

Other Information

Store at room temperature.

D&C Green no. 5, D&C yellow no.10, FD&C blue no.1, isopropyl alcohol, polyethylene glycol 300, purified water

Questions 800.835.2263

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19-001 misc	FIRST AID ONLY.
Sting F	Sting Relief Swabs
Relic	10 Swabs



Manufactured for:
Acme United Corporation
55 Walls Dr. Fairfield, CT 06824
www.FirstAidOnly.com
Made in USA
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BOX19001-revD

BOX19001-revD (002).jpg

FIRST AID ONLY STING RELIEF

benzocaine swab

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0924-5250(NDC:46414-2040)		
Route of Administration	TOPICAL				

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZO CAINE (UNII: U3RS Y48 JW5) (BENZO CAINE - UNII:U3RS Y48 JW5)	BENZOCAINE	0.12 g in 0.6 mL	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.006 g in 0.6 mL	

Inactive Ingredients				
Ingredient Name	Strength			
ISOPROPYL ALCOHOL (UNII: ND2M416302)				
WATER (UNII: 059QF0KO0R)				
D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)				
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
POLYETHYLENE GLYCOL 300 (UNII: 5655G9 Y8 AQ)				

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:0924-5250-01	10 in 1 CARTON	0 2/0 1/20 10	
1	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		0 2/0 1/20 10	

Labeler - Acme United Corporation (001180207)

Registrant - Acme United Corporation (001180207)

Establishment			
Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	relabel(0924-5250), repack(0924-5250)

Establishment				
Name	Address	ID/FEI	Business Operations	
Acme United Corporation		080119599	relabel(0924-5250), repack(0924-5250)	

Revised: 4/2018 Acme United Corporation