

STING RELIEF- alcohol, lidocaine hydrochloride cloth
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.

Drug Facts

Active Ingredient:

Ethyl Alcohol 50.0%

Lidocaine HCl 2.0%

Purpose:

Antiseptic

Analgesic (pain reliever)

Uses

First aid analgesic to help prevent infection in and provide temporary relief of the pain of:

- Insect bites and stings
- Minor scrapes and burns

Warnings:

For external use only. If swallowed, contact a doctor or poison control center immediately.

Keep out of the reach of children. Flammable, keep away from open flame.

DO NOT USE:

- In the eyes
- If you are allergic to any of the ingredients
- On raw surfaces, blistered areas, or over large areas of the body

WHEN USING THIS PRODUCT:

- Avoid contact with eyes or mucous membranes

STOP USE AND ASK A DOCTOR IF:

- Conditions worsen clear up and then recur
- The condition persists for more than 7 days
- A rash, allergic reaction, swelling, irritation or infection occur

Directions:

Packets are not child resistant. Tamper evident packaging, do not use if packet is opened or torn.

**Adults and children:
over 2 years of age:**

Remove stinger carefully, if possible. Remove wipe from packet, clean affected area thoroughly no more than 3 times daily with saturated applicator.

Children under 2:

Consult your doctor.

Other information:

Store at room temperature.

Inactive ingredients:

Benzalkonium Chloride, Menthol, Purified Water

Questions about this product:

Call 800-835-2263 with any questions about this product.

Principal Display Panel - Carton Label

19-002

MISC

FIRST AID ONLY®

Sting Relief Wipes

10 Packets

Toallitas para aliviar las molestias de picaduras

10 sobres



Principal Display Panel - Carton Label

G326

MISC

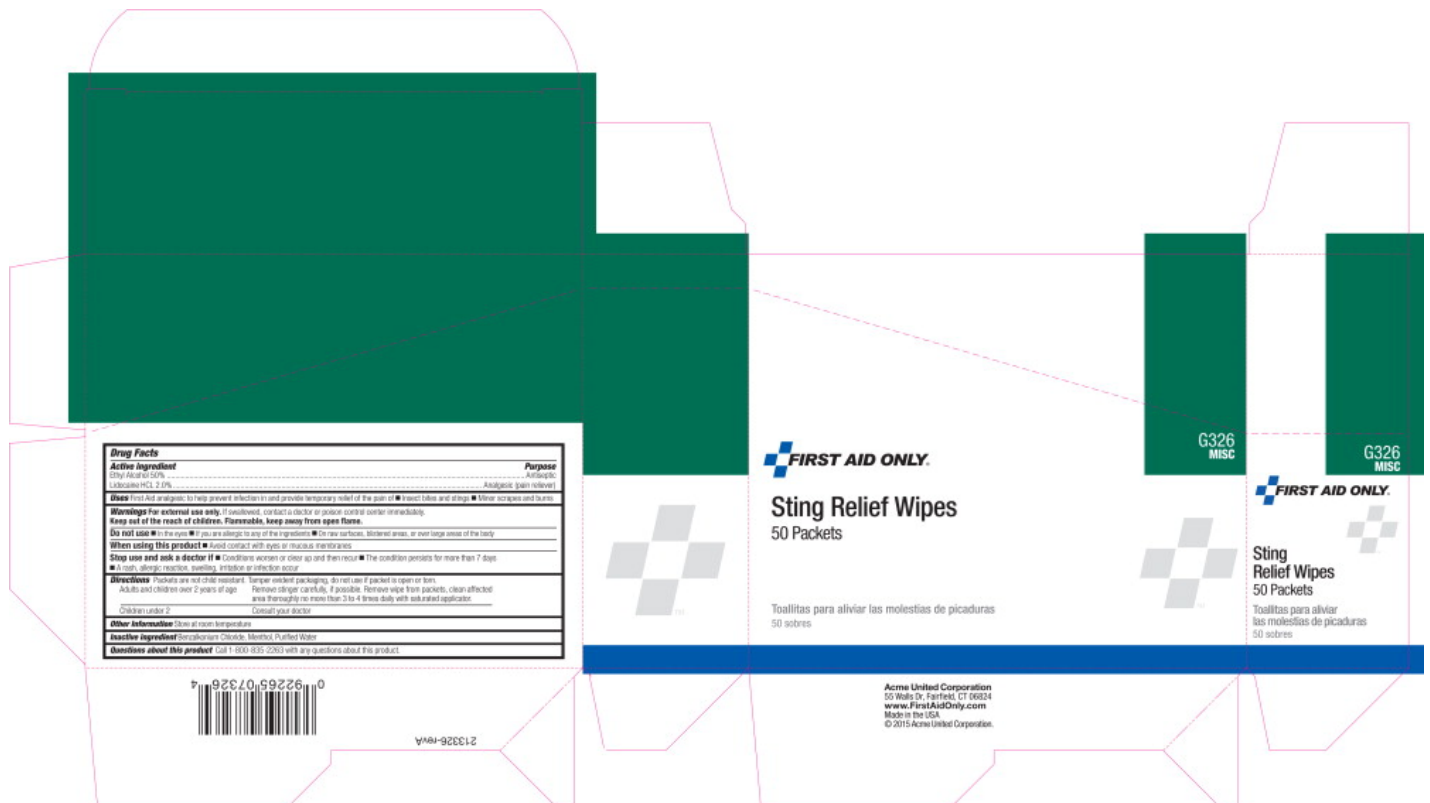
FIRST AID ONLY®

Sting Relief Wipes

50 Packets

Toallitas para aliviar las molestias de picaduras

50 sobres



Principal Display Panel - Packet Label

Sting Relief

Insect Bite Antiseptic
and Pain Reliever

1 Premoistened Towlette

Package Not Child-Resistant

FIRST AID ONLY®

www.FirstAidOnly.com

Fairfield, CT 06824

1.800.835.2263

© Acme United Corporation.

810001-revA

Sting Relief

Insect Bite Antiseptic
and Pain Reliever

1 Premoistened Towelette

Package Not Child-Resistant



www.FirstAidOnly.com
Fairfield, CT 06824
1.800.835.2263

© 2015 Acme United Corporation.
810001-revA

Drug Facts

Active Ingredients..... **Purpose**
Ethyl alcohol 50.0%..... First Aid Antiseptic
Lidocaine HCl 2.0%..... Topical Analgesic

Uses First aid to help prevent infection in minor scrapes and temporary relief of itching of insect bites.

Warnings ■ **For external use only**
■ **Flammable, keep away from fire or flame.**

Do not use ■ over large areas of the body ■ in eyes
■ over raw or blistered areas

Stop use and ask a doctor if conditions worsen or persist for more than 7 days or clear up and occur again within a few days.

Keep out of reach of children. If swallowed get medical help or contact Poison Control Center right away.

Directions Adults and childrens 2 years and older:
Apply to cleaned area not more than 3 times daily.
Children under 2 years of age: Consult a doctor.

Inactive Ingredients benzalkonium chloride,
menthol, purified water

STING RELIEF

alcohol, lidocaine hydrochloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0924-5200
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
alcohol (UNII: 3K9958V90M) (alcohol - UNII:3K9958V90M)	alcohol	0.50 mL in 1 mL
lidocaine hydrochloride (UNII: V13007Z41A) (lidocaine - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
benzalkonium chloride (UNII: F5UM2KM3W7)	
menthol (UNII: L7T10EIP3A)	
water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-5200-02	10 in 1 BOX	03/06/2012	
1	NDC:0924-5200-01	0.9 mL in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:0924-5200-03	50 in 1 BOX	03/06/2012	
2	NDC:0924-5200-01	0.9 mL in 1 PACKET; 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	part333E	03/06/2012	

Labeler - Acme United Corporation (001180207)

Registrant - Safetec of America, Inc. (874965262)

Establishment

Name	Address	ID/FEI	Business Operations
Safetec of America, Inc.		874965262	MANUFACTURE(0924-5200)

Revised: 2/2016

Acme United Corporation