BITE AND STING RELIEF WIPE- alcohol 50% lidocaine hcl 2% cloth
CVS

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.

Active Ingredients

Ethyl Alcohol 50.0%

Lidocaine HCL 2.0%

Purpose

First Aid Antiseptic
Topical Analgesic

Uses

- First aid to help guard against skin infection and temporary relief of pain and itching assocaiated with insect bites.

Warnings

For external use only.

Flammable, keep away from fire or flame.

Do not use

- **in the eyes over large areas of the body or bandage.**
- **in large quantities particularly over raw surfaces or blistered areas.**

Stop use and contact a doctor if the condition persists or worsens for more than 7 days or clears up and occurs again within a few days.

Keep out of reach of children

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

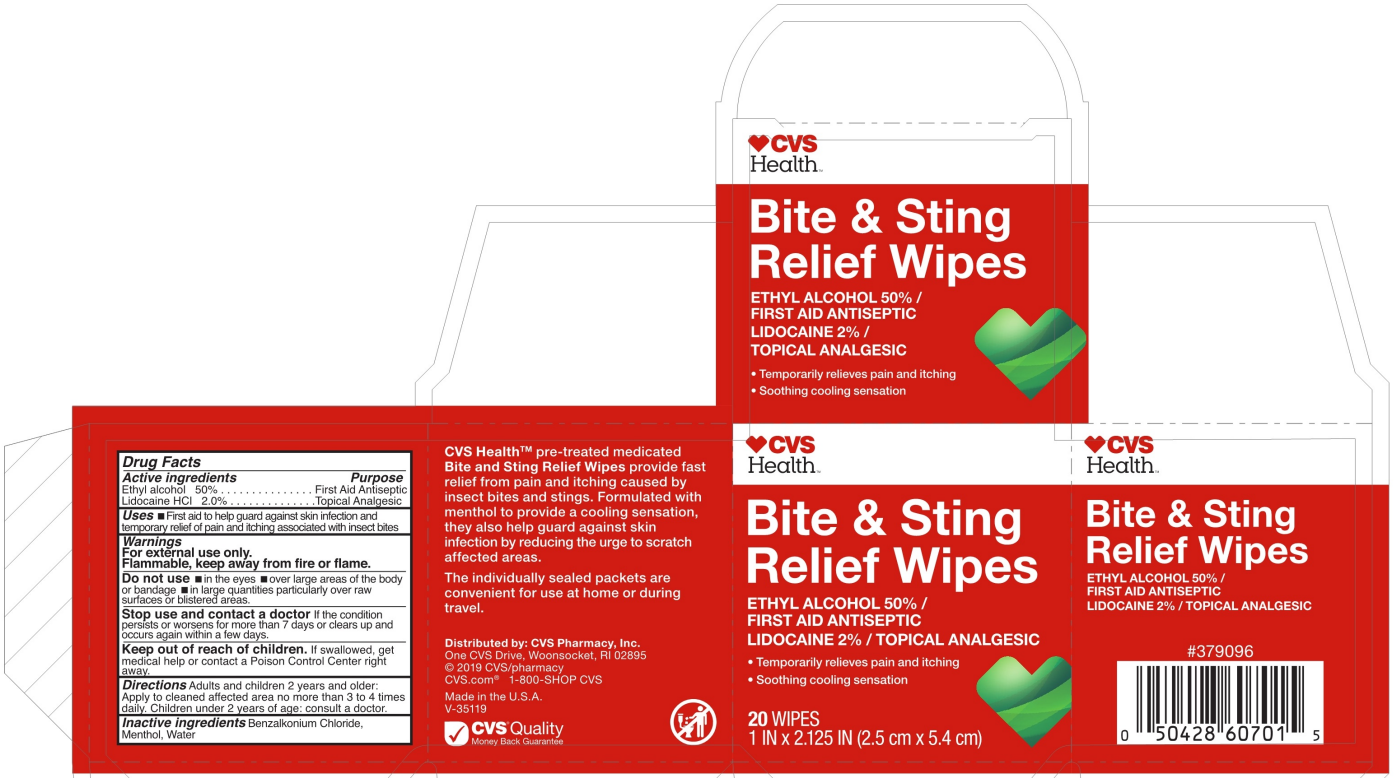
Directions

Adults and children 2 years and older: Apply to clean affected area no more than 3 to 4 times daily.
Children under 2 years of age: consult a doctor.

Inactive ingredients

Benzalkonium Chloride, Menthol, Water.

Packaging



BITE AND STING RELIEF WIPE

alcohol 50% lidocaine hcl 2% cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69842-190
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	50 mL in 100 mL
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	2 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
MENTHOL (UNII: L7T10EIP3A)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics			
Color	white	**Score**	
Shape	RECTANGLE	**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69842-190-10	30 in 1 BOX	06/01/2019	
1		3 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	part348	06/01/2019	

Labeler - CVS (062312574)

Establishment

Name	Address	ID/FEI	Business Operations
Diamond Wipes International, Inc.		161104729	manufacture(69842-190)

Revised: 7/2019 CVS