AFTER BITE WIPE INSECT STING RELIEF- benzocaine,alcohol swab
Tender 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.

After Bite Wipe Insect Sting Relief

Active Ingredients

Benzocaine, 6%

SD alcohol, 60%

Purpose

Topical Analgesic
Antiseptic

Use

temporarily relieves pain and itching due to minor stings and insect bites

Warnings

For external use only.

Flammable, keep away from fire or flame.

Do not use

- over large areas of the body
- in the eyes
- on broken skin or deep puncture wounds

Stop use and ask a doctor if

condition worsens or if symptoms persist for more than 7 days or clears up and occurs again within a few days.

Keep out of reach of children

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

Directions

- adults and children 2 year of age and older: apply to affected area not more than 3 to 4 times daily.
- children under 2 years of age: consult a doctor

Other information

Store at room temperature

Inactive Ingredient

Purified water

After Bite Wipe Insect Sting Relief

1 single use wipes

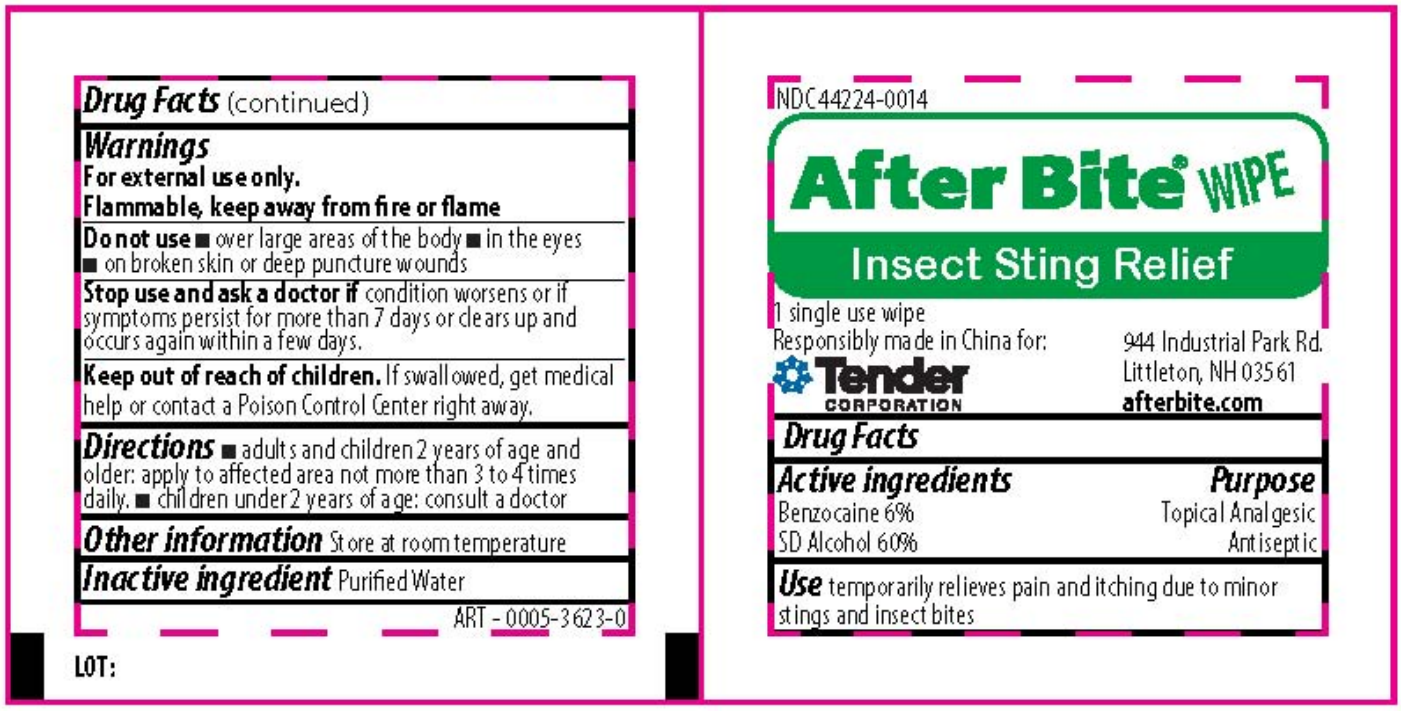
Responsibly made in China for:

Tender Corporation

944 Industrial park Rd.

Littleton, NH

afterbite.com



AFTER BITE WIPE INSECT STING RELIEF

benzocaine,alcohol swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:44224-0014
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	6 mg in 100 mg
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	60 mg in 100 mg

Inactive Ingredients	
Ingredient Name	**Strength**

WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44224-0014-1	1000 mg in 1 PACKAGE; Type 0: Not a Combination Product	09/01/2019	
2	NDC:44224-0014-2	100 in 1 BOX	09/01/2019	
2		1000 mg 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	09/01/2019	

Labeler - Tender Corporation (064437304)

Revised: 9/2019 Tender Corporation