

## **AFTER BITE WIPE- ethyl alcohol, lidocaine hcl liquid**

### **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.*

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### **After Bite Wipe**

#### **Active Ingredients**

Ethyl Alcohol 50.0%

Lidocaine HCl 2.0%

#### **Purpose**

First Aid Antiseptic

Topical Analgesic

#### **Uses**

First aid to help prevent infection in minor scrapes and temporary relief of itching 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 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 Children 2 years and older: Apply to cleaned affected area not more than 3 times daily.

Children under 2 years of age: consult a doctor.

#### **Inactive Ingredients**

benzalkonium chloride, menthol, purified water

## After Bite

After Bite

Afterbite.com

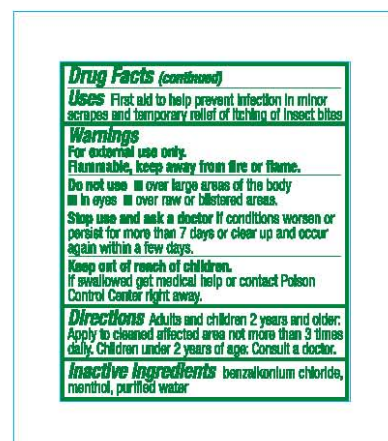
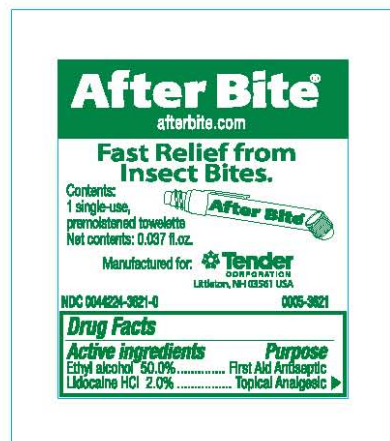
Fast Relief from Insect Bites.

Contents: 1 single-use, premoistened towelette

Net contents: 0.037fl.oz.

Manufactured for: Tender Corporation Littleton, NH 03561 USA

1.5 inches wide 1.75 inches tall



## AFTER BITE WIPE

ethyl alcohol, lidocaine hcl liquid

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44224-3621
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 mL
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.5 mL

ALCOHOL (UNII: 5K9558 V501M) (ALCOHOL - UNII:5K9558 V501M)

ALCOHOL

in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
MENTHOL (UNII: L7T10EP3A)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
WATER (UNII: 059QF0K00R)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44224-3621-0	1.1 mL in 1 POUCH; Type 0: Not a Combination Product	11/01/2016	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	11/01/2016	

**Labeler** - Tender Corporation (064437304)**Registrant** - Tender Corporation (064437304)**Establishment**

Name	Address	ID/FEI	Business Operations
Tender Corporation		064437304	manufacture(44224-3621)

Revised: 11/2016

Tender Corporation