AFTER BITE WIPE- ethyl alcohol, lidocaine hcl liquid
Adventure Ready Brands

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

Active Ingredients

Ethyl Alcohol 50.0%

Lidocaine HCl 2.0%

Purpose

First Aid Antiseptic

Topical Analgesic

Uses

First aid to help prevent infrection 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 kep 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

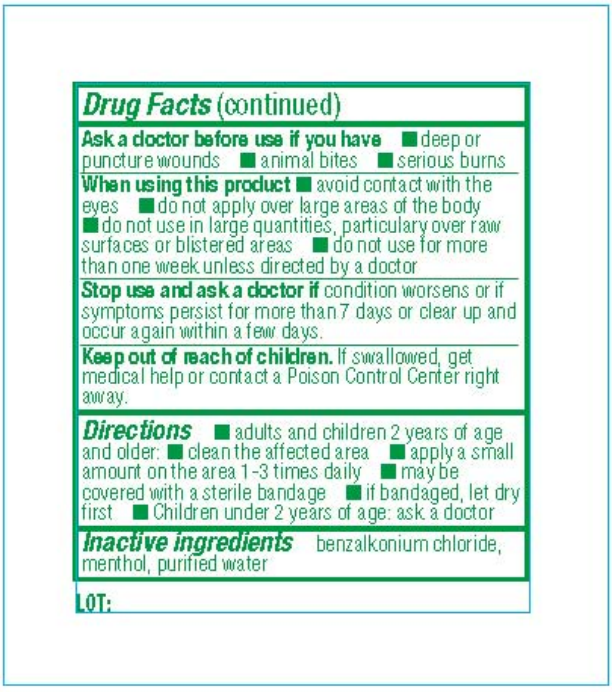
Afterbite.com

Fast Relief from Insect Bites.

Contents: 1 single-use, premoistened towelette

Net contents: 0.037fl.oz.

Manufactured for: Tender Corporation Littleton, NH 03561 USA



AFTER BITE WIPE

ethyl alcohol, lidocaine hcl liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:90107-3621
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 in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
MENTHOL (UNII: L7T10EIP3A)	

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:90107-3621-0	1.1 mL in 1 POUCH; Type 0: Not a Combination Product	01/01/2021	
2	NDC:90107-3621-1	4 in 1 BOX	01/01/2021	
2		1.1 mL in 1 POUCH; 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	01/01/2021	

Labeler - Adventure Ready Brands (064437304)

Registrant - Adventure Ready Brands (064437304)

Establishment

Name	Address	ID/FEI	Business Operations
Adventure Ready Brands		064437304	manufacture(90107-3621)

Revised: 1/2021 Adventure Ready Brands