

**BENZOCAINE,ISOPROPYL ALCOHOL- benzocaine,isopropyl alcohol swab
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 Sting Relief Pad

Drug Facts

Active Ingredient

Benzocaine, 6% w/v

Isopropyl Alcohol 60% w/v

Purpose

Topical Anesthetic

Antiseptic

Use

For temporary relief of pain and itching associated with minor burns, scrapes and insect bites.

Warnings

For external use only.

Do not use

- in the eyes If contact occurs, flush eyes with water.

Flammable,

keep away from fire or flame.

Stop use and ask a doctor if

irritation and redness develop

Keep out of reach of children

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

Directions

Apply to affected area 3-4 times daily. For adults and children 2 years of age and older. Children under 2 years; consult physician.

Inactive Ingredients

purified water

REF 856 NDC 44224-3623-0

After Bite[®] WIPE

STING RELIEF PAD

Topical Anesthetic/Antiseptic
For temporary relief of pain and itching associated with minor burns, scrapes and insect bites.
For External Use Only

Manufactured for:

 **Adventure Ready Brands™**
Made in China

1
/Pouch

 

944 Industrial Park Rd.
Littleton, NH 03561
afterbite.com

0005-3623-D
D07110807 Rev6

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BENZOCAINE, ISOPROPYL ALCOHOL

benzocaine, isopropyl alcohol swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44224-3623
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	60 mg in 1 mL
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.6 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44224-3623-0	0.4 mL in 1 POUCH; Type 0: Not a Combination Product	01/03/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	01/03/2022	

Labeler - Adventure Ready Brands (064437304)

Registrant - Adventure Ready Brands (064437304)

Establishment

Name	Address	ID/FEI	Business Operations
Adventure Ready Brands		064437304	manufacture(44224-3623)

Revised: 12/2021

Adventure Ready Brands