

**MEDICAINE STING AND BITE- benzocaine swab**  
**James Alexander Corporation**

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**MEDICAINE STING® & BITE SWABS**

***Drug Facts***

***Active Ingredients (each swab)***

Benzocaine USP (20%), L-Menthol USP (1%)

***Purpose***

analgesic

***Uses***

For temporary relief of pain and itching associated with insect bites and stings.

***Warnings***

**For external use only**

- do not use in the eyes or mouth
- not for prolonged use
- do not apply other medications to the same affected areas unless advised by a doctor

**Stop use and ask a doctor if** condition for this preparation is used persists, or if a rash or irritation or allergic reaction develops.

**Keep out of reach of children. Not for use with children less than 2 years old without medical advice.** *If swallowed, get medical help immediately or contact a Poison Control Center right away.*

***Directions***

Reverse cardboard sleeve then crush at dot between thumb and forefinger. Once solution has saturated tip, apply topically to the sting or bite. May be used on affected area(s) up to 4 times per day.

***Other Information***

Store at room temperature away from light.

***Inactive Ingredients***

D&C Green Dye #5, D&C Yellow Dye #10, FD&C Blue Dye #1, PEG 300 NF, Isopropyl Alcohol USP, Purified Water USP.

## Questions?

Call **1-908-362-9266** Monday through Friday, 9:00am - 5:00pm e.s.t

## DISPENSING SOLUTIONS®

### JAMES ALEXANDER CORPORATION

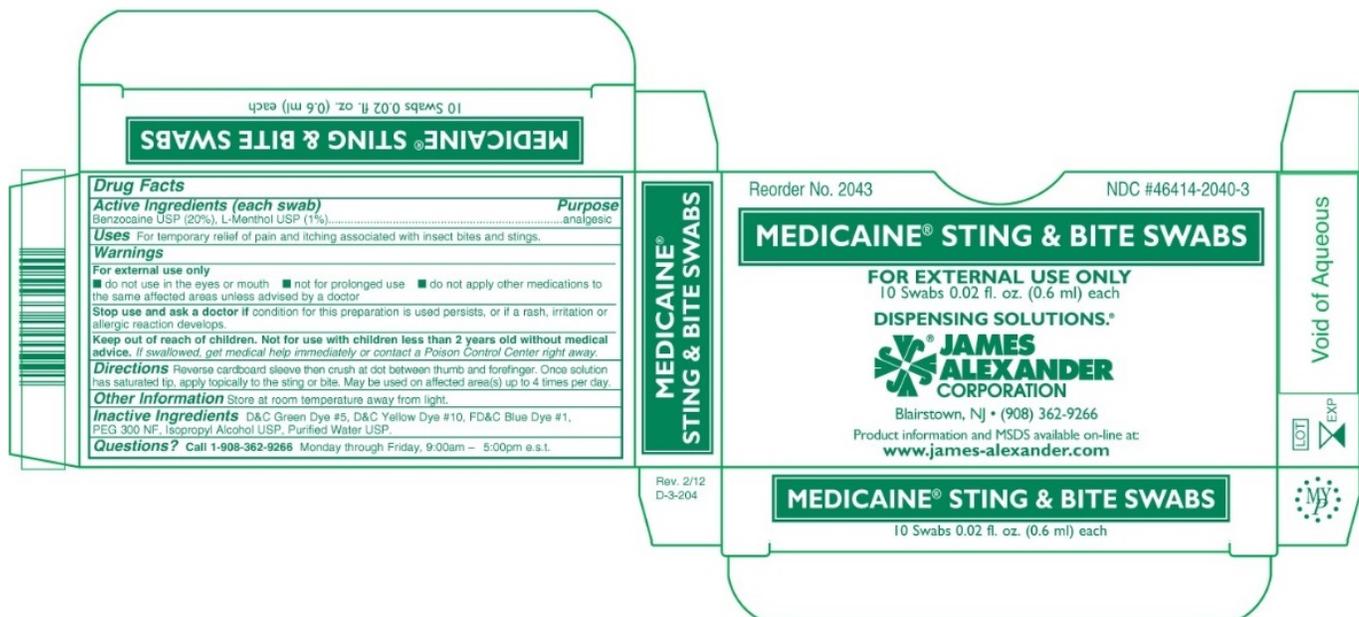
Blairstown, NJ. (908) 362-9266

Product information and MSDS available on-line at:

**www.james-alexander.com**

Void of Aqueous

## Packaging



## MEDICAINE STING AND BITE

benzocaine swab

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:46414-2040
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>BENZOCAINE</b> (UNII: U3RSY48JW5) (Benzocaine - UNII:U3RSY48JW5)	BENZOCAINE	0.12 g in 0.6 mL
<b>MENTHOL, UNSPECIFIED FORM</b> (UNII: L7T10EIP3A) (Menthol - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	0.006 g in 0.6 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>D&amp;C GREEN NO. 5</b> (UNII: 8J6RDU8L9X)	
<b>D&amp;C YELLOW NO. 10</b> (UNII: 35SW5USQ3G)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>POLYETHYLENE GLYCOL 300</b> (UNII: 5655G9Y8AQ)	
<b>ISOPROPYL ALCOHOL</b> (UNII: ND2M416302)	
<b>WATER</b> (UNII: 059QF0KOOR)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46414-2040-3	6 mL in 1 CARTON; Type 0: Not a Combination Product	11/15/1986	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/15/1986	

**Labeler** - James Alexander Corporation (040756421)**Registrant** - James Alexander Corporation (040756421)**Establishment**

Name	Address	ID/FEI	Business Operations
James Alexander Corporation		040756421	manufacture(46414-2040)

Revised: 12/2024

James Alexander Corporation