

STING RELIEF SWAB- benzocaine swab
Epic Medical Supply Corp

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

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
- 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, 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.

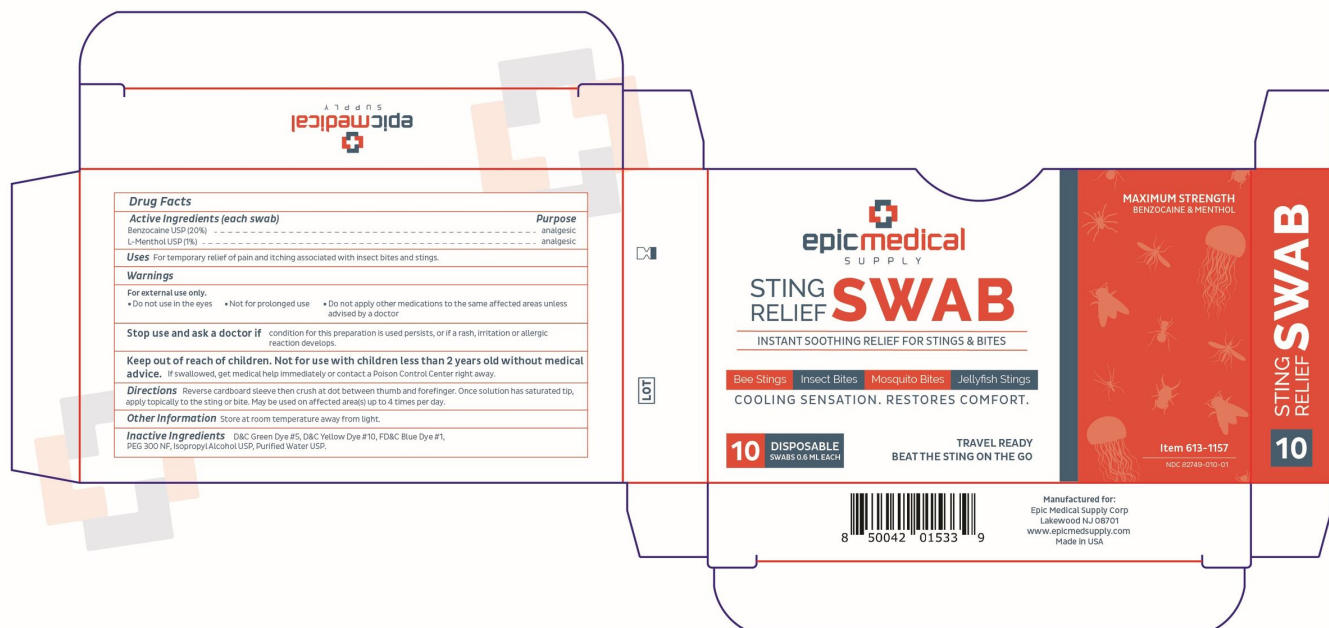
Other Information

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.

Packaging



STING RELIEF SWAB

benzocaine swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82749-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	0.006 g in 0.6 mL
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	0.12 g in 0.6 mL

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL 300 (UNII: 5655G9Y8AQ)	
WATER (UNII: 059QF0KO0R)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	

D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82749-010-01	6 mL in 1 CARTON; Type 0: Not a Combination Product	10/29/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/29/2024	

Labeler - Epic Medical Supply Corp (101423894)

Revised: 10/2024

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