MEDICAINE STING AND BITE- benzocaine swab
Dynarex Corporation

1407 Medicaine Insect Bite (Ampule) 67777-405-01

Active Ingredient

Benzocaine USP 20%

Purpose

Analgesic

Active Ingredient

L-Menthol USP 1%

Purpose

Analgesic

Use(s)

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

Warnings

For External Use Only

When Using Section

• Do not use in the eyes

• Not for prolonged use

• Do not apply other medication to the same affected areas unless advised by a doctor

Stop use and ask a doctor if

Condition persists, or if a rash, irritation, or allergic reaction develops.

Keep out of reach of children

If swallowed get medical help or contact a Poison Control Center (1-800-222-1222) 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 bit. May be used on affected area(s) up to 4 times per day

Not for use with children less than 2 years old without medical advise.

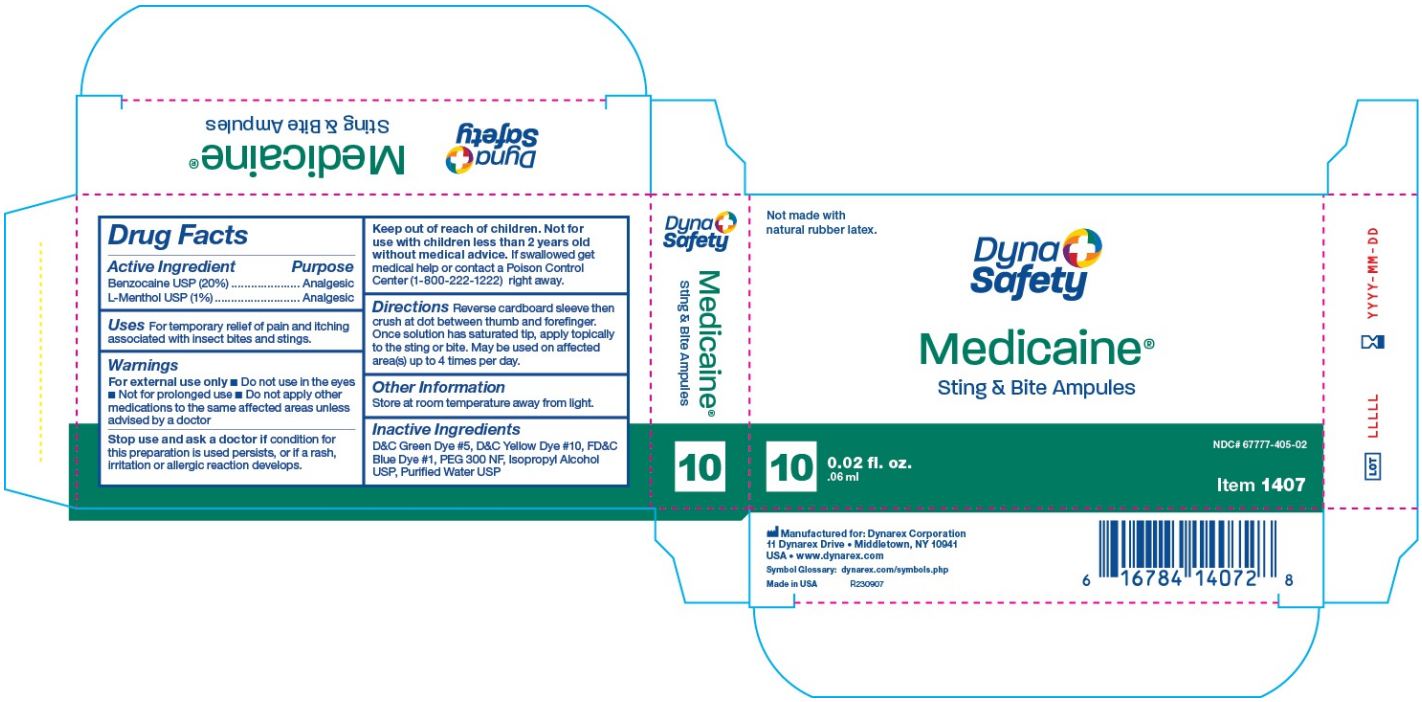
Other Information

Store at room temperature 15º-30ºC (59º-86ºF) away from light.

Inactive Ingredients

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

Label



1407 Medicaine Sting and Bite Ampule

MEDICAINE STING AND BITE

benzocaine swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-405
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	0.12 g in 0.6 mL

Inactive Ingredients

Ingredient Name	Strength
MENTHOL (UNII: L7T10EIP3A)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-405-03	50 in 1 CASE	02/14/1976	
1	NDC:67777-405-02	10 in 1 BOX		
1	NDC:67777-405-01	6 mL in 1 AMPULE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	02/14/1976	

Labeler - Dynarex Corporation (008124539)

Registrant - Dynarex Corporation (008124529)

Revised: 6/2024 Dynarex Corporation