BENZOCAINE- benzocaine swab
Dynarex Corporation

1408 Sting & Bite Pad 67777-246-01

Active Ingredient

Benzocaine, 6% w/v

Purpose

Topical Analgesic

Active Ingredient

Isopropyl Alcohol, 60% w/v

Purpose

Antiseptic

Use(s)

For the temporary relief of pain and/or itching associated with minor burns, sunburn, minor cuts, scrapes, insect bites, or minor skin irritations.

Warnings

For External Use Only

- **Flammable,** keep away from fire or flame
- Avoid contact with eyes; if this occures, rinse thoroughly with water
- Do not use with electrocautery procedures

Stop use if

- Irritation and redness develop
- Condition worsens or symptoms persists for more than 7 days, or clear up and occur again within a few days, discontinue use and consult a doctor

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

- **For adults and children 2 years of age and older:** Apply to affected area not more than 3-4 times daily.
- **Children under 2 years:** Consult a physician before use
- **For other uses:** Apply as needed

Other Information

• Store at room temperature between 15º-30ºC (59º-86ºF)

• Avoid excessive heat

Inactive Ingredients

Water

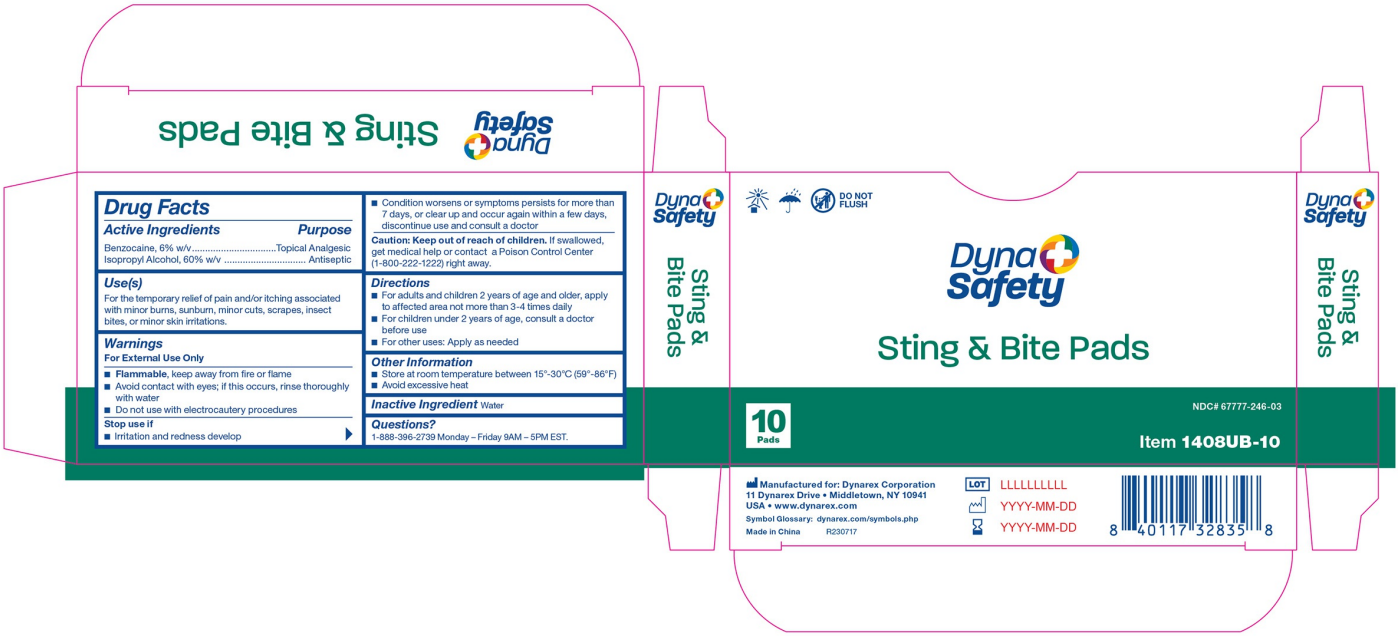
Questions?

1-888-396-2739 Monday - Friday 9AM-5PM EST.

Label

Sting & Bite Pad Label

Label 1408UB-10



1408UB-10

BENZOCAINE

benzocaine swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-246
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	60 mg in 100 mL
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	6 mg in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Product Characteristics			
Color		**Score**	
Shape	RECTANGLE	**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-246-02	3000 in 1 CASE	07/20/2011	
1	NDC:67777-246-01	2 mL in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:67777-246-04	480 in 1 CASE	07/20/2011	
2	NDC:67777-246-03	10 in 1 BOX		
2		2 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	07/20/2011	

Labeler - Dynarex Corporation (008124539)

Revised: 1/2024

Dynarex Corporation