

**BENZOCAINE- benzocaine swab**  
**Dynarex Corporation**

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**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 occurs, 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 15°-30°C (59°-86°F)
- Avoid excessive heat

## Inactive Ingredients

Water

## Questions?

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

## Label

 <p><b>Reorder No. 1408</b></p> <h1>Sting &amp; Bite Pad</h1>  <p><b>DRUG FACTS:</b></p> <table border="1"><thead><tr><th>Active Ingredients</th><th>Purpose</th></tr></thead><tbody><tr><td>Benzocaine, 6% w/v .....</td><td>Topical Analgesic</td></tr><tr><td>Isopropyl Alcohol, 60% w/v .....</td><td>Antiseptic</td></tr></tbody></table> <p><b>USE:</b> For temporary relief of pain and/or itching associated with minor burns, sunburn, minor cuts, scrapes, insect bites or minor skin irritations.</p> <p><b>WARNINGS:</b> For external use only. Flammable, keep away from fire or flame. Avoid contact with eyes; if this happens, rinse thoroughly with water. <b>DO NOT USE:</b> With electrocautery procedures.</p> 	Active Ingredients	Purpose	Benzocaine, 6% w/v .....	Topical Analgesic	Isopropyl Alcohol, 60% w/v .....	Antiseptic	<p><b>DRUG FACTS (CONTINUED)</b></p> <p><b>STOP USE:</b> If irritation and redness develop. If condition worsens or if symptoms persist for more than 7 days, or clear up and occur again within a few days, discontinue use and consult a doctor.</p> <p><b>CAUTION:</b> Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.</p> <p><b>DIRECTIONS:</b> For adults and children 2 years of age and older, apply to affected area not more than 3-4 times daily. For children under 2 years of age, consult a doctor before use.</p> <p><b>OTHER INFORMATION:</b> Store at room temperature 15°-30°C (59°-86°F). Avoid excessive heat.</p> <p><b>Inactive Ingredient:</b> Water</p> <p>Manufactured for: Dynarex Corporation 10 Glenshaw Street • Orangeburg, NY 10962 USA • www.dynarex.com</p> <p>Made in China      <b>Tear Here</b>      R190208</p>
Active Ingredients	Purpose						
Benzocaine, 6% w/v .....	Topical Analgesic						
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Sting & Bite Pad Label

## Label 1408UB-10



1408UB-10

## BENZOCAINE

benzocaine swab

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:67777-246
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ISOPROPYL ALCOHOL</b> (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	60 mg in 100 mL
<b>BENZOCAINE</b> (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	6 mg in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	

### Product Characteristics

<b>Color</b>		<b>Score</b>	
<b>Shape</b>	RECTANGLE	<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

**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: 1/2026

Dynarex Corporation