

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

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**Benzocaine Sting and Bite**

**Active Ingredient**

Active Ingredient Purpose

Benzocaine, 6% w/v                      Topical Analgesic

**Purpose for Benzocaine**

- For temporary relief of pain and itching associated with minor burns, sunburn, scrapes and insect bites or minor skin irritations

**Keep Out of Reach of Children**

- Keep out of reach of children.
- If swallowed, get medical help or contact a Poison Control Center.

**Indications and Usage**

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

**Warnings**

Warnings: Benzocaine

- For external use only.
- Flammable, keep away from fire or flame.

**Dosage**

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.

**Inactive Ingredients**

- Inactive Ingredients: Isopropyl Alcohol, Water.

**DO Not Use**

Do not use:

With electrocautery procedures.

## Stop Use

Stop use:

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.

## Storage and Handling

- Avoid excessive heat.
- Store at room temperature: 15 - 30 degrees C (59 - 86 degrees F)

## Principal Display Panel

Dynarex Sting and Bite

Benzocaine.jpg

 <p style="text-align: right;"><b>Reorder No. 1408</b> <b>NDC# 67777-246-01</b> <b>NPN #</b> <input type="text"/></p> <p style="text-align: center;"><b>Sting &amp; Bite Pad</b></p>   <p><b>DRUG FACTS:</b></p> <table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 50%;">Active Ingredient</th> <th style="width: 50%;">Purpose</th> </tr> </thead> <tbody> <tr> <td>Benzocaine, 6% w/v.....</td> <td>Topical Analgesic</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.</p> <p><b>DO NOT USE:</b> With electrocautery procedures. ▶</p>	Active Ingredient	Purpose	Benzocaine, 6% w/v.....	Topical Analgesic	<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: Keep out of reach of children.</b> 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 Ingredients:</b> Isopropyl Alcohol, Purified Water</p> <p>Manufactured for: Dynarex Corporation Orangeburg, NY 10962 www.dynarex.com</p> <p style="text-align: right;">Made in China</p> <p style="text-align: center;">- - - - Tear Here - - - -</p>
Active Ingredient	Purpose				
Benzocaine, 6% w/v.....	Topical Analgesic				

## BENZOCAINE

benzocaine swab

### Product Information

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

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOCAINE (BENZOCAINE)	BENZOCAINE	60 mg in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
ISOPROPYL ALCOHOL	
WATER	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-246-01	3000 in 1 CASE		
1		2 mL in 1 PACKET		

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part348	07/20/2011	

**Labeler** - Dynarex Corporation (008124539)**Registrant** - Dynarex Corporation (008124539)

Revised: 10/2014

Dynarex Corporation