### DEFEND- benzocaine gel Mydent International

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.

-----

# **Active Ingredients**

Benzocaine 200mg (in each g)

## Purpose

Oral Anesthetic

### Use

For oral mucosal use only, as directed by dentist. For the temporary relief of pain due to minor dental procedures.

## Warnings

**Methemoglobinemia warning:** Use of this product may cause methemoglobinemia, a serious condition that must be treated promptly because it reduces the amount of oxygen carried in the blood. This can occur even if you have used this product before. Stop use and seek immediate medical attention if you or a child in your care develops:

- pale, gray or blue colored skin (cyanosis)
- headache
- rapid heart rate
- shortness of breath
- dizziness or lightheadedness
- fatigue or lack of energy

**Allergy Alert:** Do not use on patients with a history of allergies to local anesthetics such as procaine, butacaine, benzocaine or other "caine" anesthetics.

## Do not use

- for more than 7 days unless directed by a physician. If sore mouth symptoms do not improve in 7 days; irritation, pain, or redness persists or worsens; or if swelling, rash or fever develops, see your physician promptly.
- for teething
- in children under 2 years of age

When using this product Avoid contact with eyes. If contact occurs, flush with water.

**Do not exceed recommended dosage.** If more than used for pain is accidentally swallowed, get medical help or contact a Poison Control Center right away.

If pregnant or breast feeding, ask a physician before use.

# Keep out of reach of children.

### Directions

- Apply only amount needed to the oral mucosa to prevent or relieve pain.
- children under 2 years of age: do not use

## Other information

Store at 59°-86°F (15°-30°C). Protect from freezing.

### Inactive ingredients

flavoring, PEG 3350, PEG 400, sodium saccharin. May contain blue #1, green #3, green #5, red #3, red #28, red #40, yellow #5, (tartrazine), yellow #6, as a color additive.

## Questions or comments?

1-800-275-0020



directed by a physician. If sore mouth symptoms do not improve in 7 days; irritation, pain, or redness persists or worsens; or if swelling, rash or fever develops, see your physician promptly. • for treething • in children under 2 years of age When using this product Avoid contact occurs; flush with water. Do not exceed recommended dosage. If more than used for pain is accidentally swallowed, get melical help or contact a poison Control Center right away. If pregnant or breast feeding, as a physician before use. Meep out of reach of children. Directions • Apply only amount needed to the oral muccas to prevent or relive pain. • children under 2 years of age: do not use Other information Store at 59° - 86° f (15 - 30° C). Protect from freezing. More from freezing. Inactive ingredients flavoring, PEG 3350, PEG 400, sodium saccharin. May contain blue #1, green #3, green #5, red #3, red #28, red #40, yellow #6, additive. 000-275-0020 39093151 Rev 07/2018
--

DEFEND					
penzocaine gel					
Product Information					
Product Type	HUMAN OTC DRUG	ltem Code	(Source)	NDC:	70721-003
Route of Administration	DENTAL				
A ative la ave dia at/A ative	Malaka				
Active Ingredient/Active	•				
Ingred	ient Name		Basis of Stren	gth	Strength
BENZOCAINE (UNII: U3RSY48JW5)	(BENZOCAINE - UNII:U3RSY	48JW5)	BENZOCAINE		200 mg in 1 g
Inactive Ingredients					
mactive mgreatents	In any dia at Name				Churcherth
	Ingredient Name				Strength
POLYETHYLENE GLYCOL 3350 (	JNII: G2M7P15E5P)				
POLYETHYLENE GLYCOL 400 (UI	NII: B697894SGQ)				
SACCHARIN SODIUM (UNII: SB8Z					

FD&C YELLOW NO.	<b>5</b> (UNII: 1753WB2F1M)					
FD&C YELLOW NO.	· /					
FD&C RED NO. 3 (UN	NII: PN2ZH5LOQY)					
FD&C RED NO. 40 (U	JNII: WZ B9127XOA)					
FD&C BLUE NO. 1 (U	JNII: H3R47K3TBD)					
FD&C GREEN NO. 3	(UNII: 3P3ONR6O1S)					
D&C RED NO. 28 (UN	NII: 767IP0Y5NH)					
D&C GREEN NO. 5 (U	JNII: 8J6RDU8L9X)					
Product Charac	teristics					
Color	pink	Score				
Shape		Size				
Flavor BUBBLE GUM		Imprint Code				
Contains						
Packaging						
# Item Code	Package Description	M	larketing Start Date	Marketing End Date		
	30 g in 1 JAR; Type 0: Not a Combination Product	07/0	01/2018			
Marketing Ir	nformation					
Marketing Category	Application Number or Monograp Citation	h	Marketing Start Date	Marketing End Date		
OTC monograph not final	part356		11/01/2014			

Labeler - Mydent International (176970747)

Revised: 1/2022

Mydent International