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



Drug Facts Active ingredients Berzocaine 200 mg (in each g) Purpose Dral Anesthetic Use For oral mucosal use only, as directed by dentist. For the temporary relief of pain due to minor dental procedures. Warnings Methemoglobinemia, a serious condition that must be treated promptly because it readuces the amount of oxygen carried in the blood. This can occur even if you have used this product before. Stop use and 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 carried the amount of oxygen seek immediate medical attention fiy ou or a child in your a cut soro relach 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

DEFEND					
oenzocaine gel					
Product Information					
Product Type	HUMAN OTC DRUG	ltem Code	(Source)	NDC:	70721-007
Route of Administration	DENTAL				
Active Ingredient/Active	Maiaty				
Active Ingredient/Active	•		Basis of Stren		- · · ·
Ingred	gth	Strength			
BENZOCAINE (UNII: U3RSY48JW5)	BENZOCAINE		200 mg in 1 g		
Inactive Ingredients					
		Strength			
POLYETHYLENE GLYCOL 3350 (JNII: G2M7P15E5P)				
POLYETHYLENE GLYCOL 400 (UI	NII: B697894SGQ)				
SACCHARIN SODIUM (UNII: SB8Z					

FD&C YELLOW NO.	5 (UNII: 175	3WB2F1M)				
FD&C YELLOW NO.	6 (UNII: H7	7VEI93A8)				
FD&C RED NO. 3 (UN	NII: PN2ZH5	SLOQY)				
FD&C RED NO. 40 (U	JNII: WZ B93	127XOA)				
FD&C BLUE NO. 1 (U	JNII: H3R47	K3TBD)				
FD&C GREEN NO. 3	(UNII: 3P3C	DNR601S)				
D&C RED NO. 28 (UN	NII: 767IP0Y	′5NH)				
D&C GREEN NO. 5 (UNII: 8J6RDI	U8L9X)				
Product Charac	toristic	e				
Color red			Scor	•		
Shape		Size				
-				nprint Code		
Contains						
Packaging						
# Item Code	Package Description			Marketing Start Date	Marketing End Date	
	30 g in 1 JA Product	R; Type 0: Not a Combination	07	/01/2018		
Marketing Ir	nforma	ition				
Marketing Category	Applic	cation Number or Monogra Citation	ph	Marketing Start Date	Marketin Date	-
OTC monograph not	part356			11/01/2014		
final						

Labeler - Mydent International (176970747)

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