

ORAJEL MEDICATED FOR TOOTHACHE AND GUM, LIQUID- benzocaine and menthol liquid
Church & Dwight Co., Inc.

Orajel Medicated for Toothache and Gum, Liquid

Benzocaine 20%

Menthol 0.1%

Oral Pain Reliever

Use for the temporary relief of pain due to • canker sores • minor injury of the mouth and gums • minor irritation of the mouth and gums caused by dentures or orthodontic appliances

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 or breath • dizziness or lightheadedness • fatigue or lack of energy

Allergy Alert: do not use this product if you have a history of allergy to local anesthetics such as procaine, butacaine, benzocaine, or other "caine" anesthetics

Flammable: keep away from fire or flame. Avoid smoking during application.

Do not use • more than directed • for more than 7 days unless directed by a dentist or doctor • for teething • in children under 2 years of age

Stop use and ask a physician if • swelling, rash or fever develops • irritation, pain or redness persists or worsens • symptoms do not improve in 7 days

Keep out of reach of children.

In case of overdose or allergic reaction, get medical help or contact a Poison Control Center right away

Directions • remove imprinted safety seal from bottle cap

Adults and children 2 years of age and over

- Apply product with cotton swab or finger to the affected area. Use up to 4 times daily or as directed by a dentist or doctor

Children under 12 years of age

- Ask a doctor before use. Should be supervised in the use of this product

Children under 2 years of age

- Do not use

Other Information • do not use if safety seal on bottle cap is broken or missing prior to opening

Inactive Ingredients alcohol (54.5% by volume), flavor, PEG-8, red 40, sodium saccharin, water, yellow 5

Questions or comment? Call us at **800 952 5080** M-F 9am-5pm or visit our website at **www.oraljel.com**

#1

ORAL PAIN

RELIEVER BRAND

FOR TOOTHACHE

Fast-Acting Liquid!

Orajel

MEDICATED

FOR TOOTHACHE & GUM

Instant Pain Relief

•20% Benzocaine to Relieve Oral Pain

•Menthol to Soothe Gums

CANKER SORES • CHEEK BITES • DENTURE PAIN

IRRITATION FROM BRACES

ORAL PAIN RELIEVER

0.45 FL OZ (13.3mL)

#1 ORAL PAIN RELIEVER BRAND FOR TOOTHACHE

Fast-Acting Liquid!

Orajel™

MEDICATED

FOR TOOTHACHE & GUM

INSTANT PAIN RELIEF



✓ 20% Benzocaine to Relieve Oral Pain
✓ Menthol to Soothe Gums

CANKER SORES • CHEEK BITES • DENTURE PAIN
IRRITATION FROM BRACES

✓ 20% Benzocaine to Relieve Oral Pain
✓ Menthol to Soothe Gums

CANKER SORES • CHEEK BITES • DENTURE PAIN
IRRITATION FROM BRACES

ORAL PAIN RELIEVER

0.45 FL OZ (13.3 mL)

Church & Dwight Co., Inc.
Zwing, NJ 08628, USA

The maker of the Orajel™ brand does not
manufacture store brand oral pain products.
JUPC-32945-01 72013333



Drug Facts

Active Ingredients
Benzocaine 20%
Menthol 1%
Oral pain reliever

Purpose
Oral pain reliever

Use
For the temporary relief of pain due to:
■ canker sores ■ minor injury of the mouth and gums
■ minor irritation of the mouth and gums caused by dentures or orthodontic appliances

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 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
■ allergic alert: do not use this product if you have a history of allergy to local anesthetics such as procaine, tetracaine, benzocaine or other "caine" anesthetics

Flammability: Keep away from fire or flame. Avoid smoking during application.

Do not use ■ more than directed ■ for more than 7 days unless directed by a dentist or doctor ■ for numbing ■ in children under 2 years of age

Stop use and ask a doctor if ■ swelling, rash or fever develops ■ irritation, pain or redness persists or worsens ■ symptoms do not improve in 7 days

Keep out of reach of children. In case of overdose or allergic reaction, get medical help or contact a Poison Control Center right away.

Directions ■ remove safety seal from bottle cap
Apply product with cotton swab or finger to the affected area.
Use up to 4 times daily or as directed by a dentist or doctor.
Children between 2 and 12 years of age
Ask a doctor before use. Should be supervised in the use.
Do not use
Children under 2 years of age

Other information ■ do not use if safety seal on bottle cap is broken or missing prior to opening

Inactive ingredients alcohol (54.5% by volume), flavor, FD&C red #40, sodium saccharin, water, yellow 5

Questions or comments? call us at 1-800-952-5080 M-F 9am-5pm ET or visit our website at www.orajel.com

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ORAJEL MEDICATED FOR TOOTHACHE AND GUM, LIQUID

benzocaine and menthol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10237-796
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	200 mg in 1 mL
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
METHYL SALICYLATE (UNII: LAV5U5022Y)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
ALCOHOL (UNII: 3K9958V90M)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	orange	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10237-796-45	1 in 1 CARTON	08/07/2020	
1		13.3 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M022	08/07/2020	

Labeler - Church & Dwight Co., Inc. (001211952)

