

# **ORAJEL 3X MEDICATED FOR TOOTHACHE AND GUM- benzocaine, menthol, zinc chloride gel**

**Church & Dwight Co., Inc.**

*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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## **Orajel 3X Medicated For Toothache and Gum, Gel**

Benzocaine 20%

Menthol 0.26%

Zinc chloride 0.15%

Oral pain reliever and Oral astringent

### **Use**

for the temporary relief of pain due to minor irritation or injury of the mouth and gums

**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

**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

- 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 doctor if**

- symptoms do not improve in 7 days
- swelling, rash or fever develops
- irritation, pain or redness persists or worsens

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

### **Directions**

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Adults and children 2 years of age and over      Apply to affected area 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 of this product |
| Children under 2 years of age          | Do not use   |

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### **Other information**

do not use if Tamper-Evident Tab is open before first use

ammonium glycyrrhizate, flavor, PEG-8, PEG-75, sodium saccharin, sorbic acid, water

### **Questions or comments?**

call us at 1-800-952-5080 M-F 9am-5pm ET or visit our website at [www.oraljel.com](http://www.oraljel.com)

#1 ORAL PAIN

RELIEF BRAND

FOR ADULTS

EXTRA  
STRENGTH

Orajel™

3X TOOTHACHE & GUM

MEDICATED

IMMEDIATE PAIN RELIEF

ORAL PAIN

RELIEVER/

ASTRINGENT

Provides

Long-Lasting

Pain Relief\*

Soothes

Irritated Gums

Delivers Targeted

Pain Relief

GEL

NET WT

0.25 OZ (7.0 g)

#1 ORAL PAIN RELIEF BRAND FOR ADULTS

EXTRA STRENGTH

**Orajel™**  
**3X TOOTHACHE & GUM MEDICATED**

IMMEDIATE PAIN RELIEF

ORAL PAIN RELIEF/ANESTHESIA

- ✓ Provides Long-Lasting Pain Relief\*
- ✓ Soothes Irritated Gums
- ✓ Delivers Targeted Pain Relief

SCAN ME



LEARN MORE

- ✓ Provides Long-Lasting Pain Relief\*
- ✓ Soothes Irritated Gums
- ✓ Delivers Targeted Pain Relief

GEL

NET WT 0.25 OZ (7.0 g)

**Orajel**

TAMPER-EVIDENT TAB

Church & Dwight Co., Inc.  
 Teaneck, NJ 07666 USA

Orajel is a trademark of Church & Dwight Co., Inc. The maker of the Orajel® brand does not manufacture other brand oral pain products.

\*Up to 2 Hours



875-5540-01 700076



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**Drug Facts**

**Active Ingredients**  
 Benzocaine 20%  
 Menthol 0.25%  
 Zinc Chloride 0.15%  
 Oral Anesthetic  
 Oral Pain Reliever

**Purpose**  
 Oral Pain Reliever

**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 the 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 this product if you have a history of allergy to local anesthetics such as procaine, tetracaine, benzocaine or other "caine" anesthetics. 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 ■ swelling, rash, or fever develops ■ irritation, pain or redness persists or worsens. 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**  
 Adults and children 2 years of age and over ■ as directed by a dentist or doctor. Apply to affected area up to 4 times daily. Children between 2 and 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 Tamper-Evident Tab is open before first use. Inactive ingredients ■ artemisinin glycololate, flavor, PEG-9, PEG-15, sodium acetate, sorbic acid, water.

**Questions or comments?** call us at 1-800-552-5000 M-F 9am-5pm ET or visit our website at [www.orajel.com](http://www.orajel.com)

**ORAJEL 3X MEDICATED FOR TOOTHACHE AND GUM**  
 benzocaine, menthol, zinc chloride gel

**Product Information**

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:10237-790 |
| <b>Route of Administration</b> | TOPICAL        |                           |               |

**Active Ingredient/Active Moiety**

| <b>Ingredient Name</b>  | <b>Basis of Strength</b> | <b>Strength</b> |
|---|--------------------------|-----------------|
| <b>MENTHOL</b> (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)           | MENTHOL                  | 2.6 mg in 1 g   |
| <b>BENZOCAINE</b> (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)     | BENZOCAINE               | 200 mg in 1 g   |
| <b>ZINC CHLORIDE</b> (UNII: 86Q357L16B) (ZINC CATION - UNII:13S1S8SF37) | ZINC CATION              | 1.5 mg in 1 g   |

**Inactive Ingredients**

| <b>Ingredient Name</b>                            | <b>Strength</b> |
|---|-----------------|
| <b>WATER</b> (UNII: 059QF0KO0R)                   |                 |
| <b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ) |                 |
| <b>METHYL SALICYLATE</b> (UNII: LAV5U5022Y)       |                 |
| <b>SACCHARIN SODIUM</b> (UNII: SB8ZUX40TY)        |                 |
| <b>AMMONIUM GLYCYRRHIZATE</b> (UNII: 3VRD35U26C)  |                 |
| <b>PEG-75 LANOLIN</b> (UNII: 09179OX7TB)          |                 |
| <b>SORBIC ACID</b> (UNII: X045WJ989B)             |                 |

**Packaging**

| <b>#</b> | <b>Item Code</b> | <b>Package Description</b>                         | <b>Marketing Start Date</b> | <b>Marketing End Date</b> |
|----------|------------------|--|-----------------------------|---------------------------|
| 1        | NDC:10237-790-25 | 1 in 1 PACKAGE                                     | 11/01/2019                  |                           |
| 1        |                  | 2.5 g in 1 TUBE; Type 0: Not a Combination Product |                             |                           |
| 2        | NDC:10237-790-42 | 1 in 1 PACKAGE                                     | 11/01/2019                  |                           |
| 2        |                  | 4.2 g in 1 TUBE; Type 0: Not a Combination Product |                             |                           |
| 3        | NDC:10237-790-12 | 1 in 1 PACKAGE                                     | 11/01/2019                  | 06/10/2023                |
| 3        |                  | 3.5 g in 1 TUBE; Type 0: Not a Combination Product |                             |                           |

**Marketing Information**

| <b>Marketing Category</b> | <b>Application Number or Monograph Citation</b> | <b>Marketing Start Date</b> | <b>Marketing End Date</b> |
|---------------------------|---|-----------------------------|---------------------------|
| OTC monograph not final   | part348   | 11/01/2019                  |                           |

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

## Establishment

| Name                      | Address | ID/FEI    | Business Operations    |
|---------------------------|---------|-----------|------------------------|
| Church & Dwight Co., Inc. |         | 043690812 | manufacture(10237-790) |

Revised: 1/2022

Church & Dwight Co., Inc.