ORA YOGU 4X MEDICATED TOOTH AND GUM PAIN RELIEVER- medicated tooth and gum pain reliever liquid
Kungpo Health Care & Herbal Center Inc

DRUG FACTS

Drug Facts

Active ingredients

Benzalkonium Chloride 0.13%
Benzocaine 20%
Menthol 0.26%
Zinc Chloride 0.15%

Purpose

Benzalkonium Chloride Oral Antibacterial
Benzocaine Oral Antibacterial
Menthol Oral Antibacterial
Zinc Chloride Oral Antibacterial

Uses

- Temporarily relieves pain caused by minor oral and gum injuries.
- First aid to help prevent infection from minor oral irritations.
- Improves loose teeth and maintains dental health.

Warning

Use of this product may cause methemoglobinemia, a serious condition that requires immediate treatment because it reduces the oxygen-carrying capacity of the blood. This can happen even if you have used this product before.

Do not use

- Do not use this product if you have a history of allergy to procaine, butacaine, benzocaine, or other "caine" class local anesthetics.
- Do not exceed the recommended dosage for more than 7 days unless directed by a dentist or doctor.

Stop use and ask a doctor if

- Stop use and seek immediate medical attention if you or 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
- Sore mouth symptoms do not improve in 7 days.
- 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

- Adult 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 using this product.
- Children under 2 years of age - Do not use.

Other information

- This preparation is intended for use in cases of toothache, only as a temporary expedient until a dentist can be consulted. Do not use continuously.
- Do not use if temper-evident tab is open before first use.

Inactive ingredients

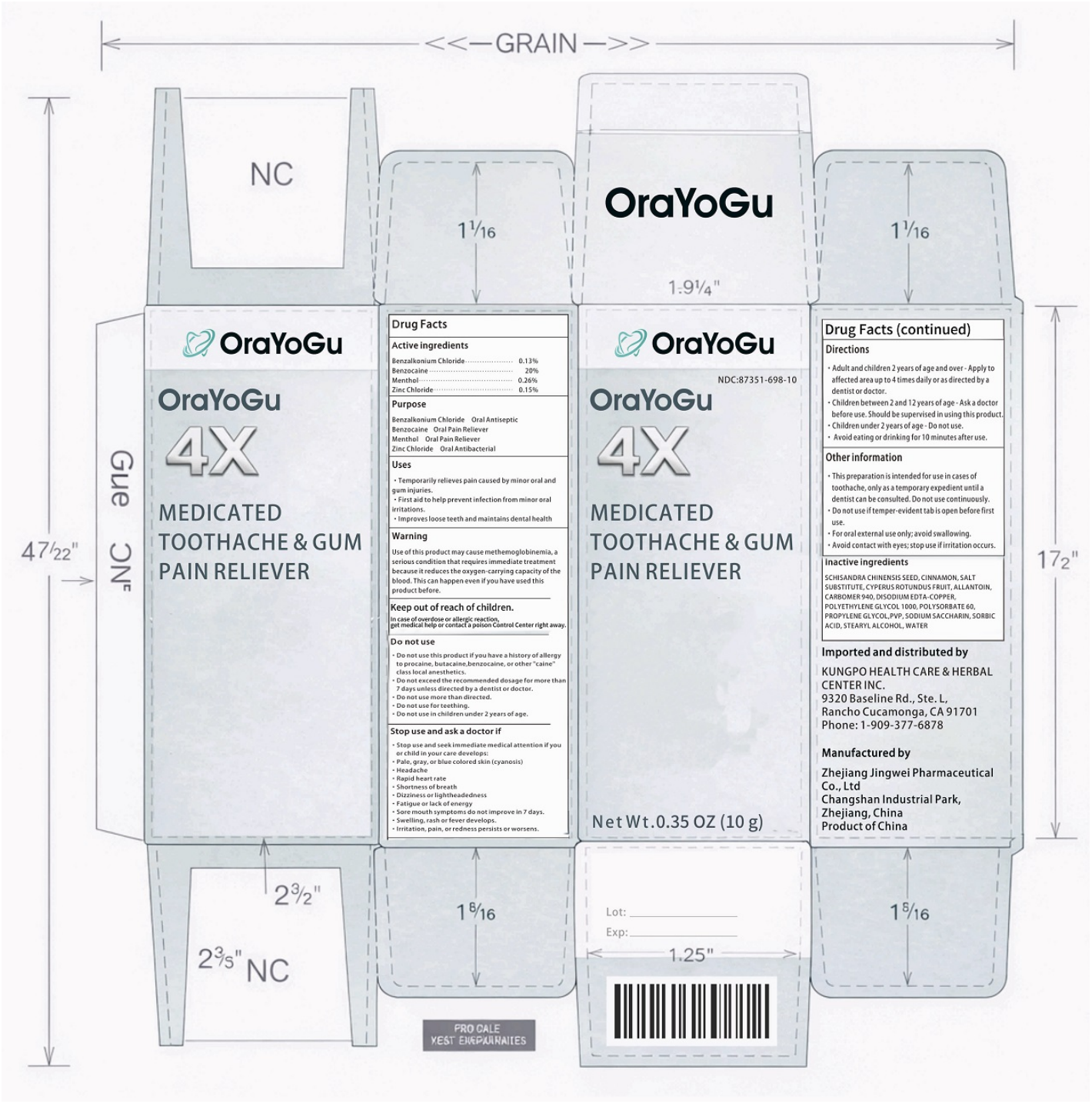
SCHISANDRA CHINENSIS SEED, CINNAMON, SALT SUBSTITUTE, CYPERUS ROTUNDUS FRUIT, ALLANTOIN, CARBOMER 940, DISODIUM EDTA-COPPER, POLYETHYLENE GLYCOL 1000, POLYSORBATE 60, PROPYLENE GLYCOL, PVP, SODIUM SACCHARIN, SORBIC ACID, STEARYL ALCOHOL, WATER

Manufactured by

Zhejiang Jingwei Pharmaceutical Co., Ltd
Changshan Industrial Park, Zhejiang, China
Product of China

Imported and distributed by

KUNGPO HEALTH CARE & HERBAL CENTER INC.
9320 Baseline Rd., Ste. L
Rancho Cucamonga, CA 91701
Phone: 909-377-6878



ORA YOGU 4X MEDICATED TOOTH AND GUM PAIN RELIEVER

medicated tooth and gum pain reliever liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:87351-698
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	20 g in 100 mL
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.26 g in 100 mL
ZINC CHLORIDE (UNII: 86Q357L16B) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	0.15 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SCHISANDRA CHINENSIS SEED (UNII: MO9DK65G18)	
CINNAMON (UNII: 5S29HWU6QB)	
SALT SUBSTITUTE (UNII: HJT620308C)	
CYPERUS ROTUNDUS FRUIT (UNII: VD727WZ56)	
ALLANTOIN (UNII: 344S277G0Z)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
DISODIUM EDTA-COPPER (UNII: 6V475AX06U)	
POLYETHYLENE GLYCOL 1000 (UNII: U076Q6Q621)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PVP (UNII: FZ989GH94E)	
SODIUM SACCHARIN (UNII: SB8ZUX40TY)	
SORBIC ACID (UNII: X045WJ989B)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor	MINT	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87351-698-10	1 in 1 PACKAGE	11/25/2027	
1		10 mL in 1 TUBE; 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	11/25/2025	

Labeler - Kungpo Health Care & Herbal Center Inc (118869850)

Registrant - Zhejiang Jingwei Pharmaceutical Co., Ltd (530876549)

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Jingwei Pharmaceutical Co., Ltd		530876549	manufacture(87351-698)

Revised: 1/2026 Kungpo Health Care & Herbal Center Inc