

**WALGREENS MOUTH SORE RELIEF- benzocaine liquid**  
**Walgreens**

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**5820635 Walgreens Mouth Sore Relief (Kanka, 8015303)**

Benzocaine 20%

Oral anesthetic/analgesic

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

**Keep out of reach of children.** If more than used for pain is accidentally swallowed, get medical help or contact a Poison Control Center right away.

**adults and children 2 years of age and over:** to assure formation of long-lasting film coating, dry affected area and apply medication undiluted with a cotton swab or fingertip; allow a few seconds for coating to form; use up to 4 times daily, or as directed by a dentist or doctor

**children under 12 years of age** should be supervised in the use of this product

**children under 2 years of age:** do not use

benzyl alcohol, cetylpyridinium chloride, compound benzoin tincture, dimethyl isosorbide, ethylcellulose, flavor, octylacrylamide/acrylates/butylaminoethyl/methylacrylate copolymer, oleth-10, polyethylene glycol, propylene glycol, ricinus communis (castor) seed oil, SD alcohol 38B, sucralose, tannic acid



**TAMPER-EVIDENT: DO NOT USE IF OUTER PACKAGING IS MISSING OR BROKEN**  
**RETAIN CARD FOR COMPLETE PRODUCT INFORMATION**

**Drug Facts**

**Active ingredient** Benzocaine 20.0% **Purpose** Oral anesthetic/analgesic

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

**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

**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

**Do not use** • for teething • in children under 2 years of age

**When using this product** • do not use for more than 7 days unless directed by a dentist or doctor. If sore mouth symptoms do not improve in 7 days; if irritation, pain or redness persists or worsens; if swelling, rash or fever develops, see your dentist or doctor promptly. • do not exceed recommended dosage.

**Keep out of reach of children.** If more than used for pain is accidentally swallowed, get medical help or contact a Poison Control Center right away.

**Directions** • adults and children 2 years of age and over: to assure formation of long-lasting film coating, dry affected area and apply medication undiluted with a cotton swab or fingertip; allow a few seconds for coating to form; use up to 4 times daily, or as directed by a dentist or doctor • children under 12 years of age should be supervised in the use of this product • children under 2 years of age: do not use

**Other information** • do not purchase if package has been opened • cap tightly after use to avoid evaporation • avoid contact with eyes • avoid contact with clothing and household/furniture surfaces to prevent possible staining • this is a personal care item, and should be used by one individual only

**Inactive ingredients**  
benzyl alcohol, cetylpyridinium chloride, compound benzoin tincture, dimethyl isosorbide, ethylcellulose, flavor, octylacrylamide/acrylates/butylaminoethyl/methylacrylate copolymer, oleth-10, polyethylene glycol, propylene glycol, ricinus communis (castor) seed oil, SD alcohol 38B, sucralose, tannic acid

**Recycling Information:**  
Remove From Card: PAPER CARD  
Empty & Replace Cap: PLASTIC TRAY  
Glass Bottle: GLASS BOTTLE

DISTRIBUTED BY: WALGREEN CO., DEERFIELD, IL 60015  
**100% SATISFACTION GUARANTEED**  
walgreens.com 1-800-925-4733  
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ITEM 771861 W00000-0000-0  
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2824635R1/5820635 NDC 0363-5334-00

WALGREENS MOUTH SORE RELIEF			
benzocaine liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-5334
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)		BENZOCAINE	200 mg in 1 mL
Inactive Ingredients			
Ingredient Name			Strength
OLETH-10 (UNII: JD797EF70J)			
DIMETHYL ISOSORBIDE (UNII: SA6A6V432S)			
POLYETHYLENE GLYCOL 300 (UNII: 5655G9Y8AQ)			
BENZOIN, (+/-)- (UNII: L7J6A1NE81)			
ACRYLATE/BUTYLAMINOETHYL METHACRYLATE/METHYL ACRYLATE/METHYL			

<b>METHACRYLATE/OCTYLACRYLAMIDE COPOLYMER (40000 WAMW)</b> (UNII: 8RZ43KFB5K)	
<b>CETYLPIRIDINIUM CHLORIDE</b> (UNII: D9OM4SK49P)	
<b>BENZYL ALCOHOL</b> (UNII: LKG8494WBH)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>ETHYLCELLULOSE, UNSPECIFIED</b> (UNII: 7Z8S9VYZ4B)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>RICINUS COMMUNIS SEED</b> (UNII: 7EK45FN1TX)	
<b>TANNIC ACID</b> (UNII: 28F9E0DJY6)	

### Product Characteristics

<b>Color</b>	brown (Light to dark brown)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	MINT (alcohol and mint)	<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-5334-00	1 in 1 BLISTER PACK	12/01/2024	
1		14.7 mL in 1 BOTTLE; 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	10/31/2024	

**Labeler** - Walgreens (008965063)

**Registrant** - Lornamead, Inc (080046418)

### Establishment

Name	Address	ID/FEI	Business Operations
HK Kolmar Canada, Inc		243501959	manufacture(0363-5334) , pack(0363-5334)