

MOUTH SORE RELIEF PROFESSIONAL STRENGTH- benzocaine liquid
CVS Pharmacy

5820303 CVS Mouth Sore Relief

Active ingredient

Benzocaine 20% (w/w)

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

For oral use only

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

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; or if swelling, rash or fever develops, see your doctor or dentist promptly. Do not exceed recommended dosage.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center

right away.

Directions

- to assure formation of long-lasting film coating, dry affected area and apply medication undiluted with applicator
- 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:** consult a dentist or doctor

Other Information

- do not use if package has been opened
- store at 20-25°C (68-77°F)

- close 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/methacrylate Copolymer, Oleth-10, PEG-6, Propylene Glycol, Ricinus Communis (Castor) Seed Oil, SD Alcohol 38B, Sucralose, Tannic Acid.

Drug Facts

Active ingredient Benzocaine 20.0% (w/w)	PurposeOral anesthetic/analgesic
Use 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 • more than directed • for more than 7 days unless told to do so by a dentist or doctor • for teething • 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 • allergic reaction occurs	
Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.	
Directions • to assure formation of long-lasting film coating, dry affected area and apply medication undiluted with applicator • 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 • store at 20-25°C (68-77°F) • close 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 methacrylate copolymer, oleth-10, PEG-6, propylene glycol, ricinus communis (castor) seed oil, SD alcohol 38B, sucralose, tannic acid.	Distributed by: CVS Pharmacy, Inc. One CVS Drive, Woonsocket, RI 02895 © 2019 CVS/pharmacy CVS.com® 1-800-SHOP CVS Made in Canada V-14931

CVS Quality
Money Back Guarantee

#711642

2824303R8/5820303 0 50428 33568 0

*This product is not manufactured or distributed by Blistex, Inc., owner of the registered trademark Blistex® and Kank-A®. Retain card for complete product information.

MOUTH SORE RELIEF PROFESSIONAL STRENGTH

benzocaine liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59779-830
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 g

Inactive Ingredients	
Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
ETHYLCELLULOSES (UNII: 7Z8S9VYZ4B)	
TANNIC ACID (UNII: 28F9E0DJY6)	
CETYLPYRIDINIUM CHLORIDE (UNII: D9OM4SK49P)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
OLETH-10 (UNII: JD797EF70J)	
DIMETHYL ISOSORBIDE (UNII: SA6A6V432S)	
RICINUS COMMUNIS SEED (UNII: 7EK4SFN1TX)	
POLYETHYLENE GLYCOL 300 (UNII: 5655G9Y8AQ)	
BENZOIN, (+/-) (UNII: L7J6A1NE81)	

Product Characteristics			
Color		Score	
Shape		Size	
Flavor	MINT (Mint Flavor #25797)	Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59779-830-38	1 in 1 BLISTER PACK	04/19/2011	
1		14.7 g in 1 BOTTLE, WTH APPLICATOR; Type 0: Not a Combination Product		
2	NDC:59779-830-48	1 in 1 BLISTER PACK	04/19/2011	
2		14.7 g 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	10/01/2009	

Labeler - CVS Pharmacy (062312574)

Establishment			
Name	Address	ID/FEI	Business Operations
HK KOLMAR CANADA, INC		243501959	manufacture(59779-830) , pack(59779-830)