BVIBE SPEARMINT DEEP THROAT- benzocaine numbing spray liquid
Private Label Productions LLC

PLP-COTR Deep Throat

Active Ingredient(s)

Benzocaine 2.5%

Purpose

Oral Anesthetic

Use

Temporarily reduces gag reflex by desensitizing the throat for deeper oral activity.

Warnings

For external use only. Do not apply to genitals or other body parts.
Allergy Alert: Avoid if allergic to benzocaine or other local anesthetics (e.g., lidocaine, procaine)
Do Not Use: If you have a history of methemoglobinemia or take related medications (e.g., nitrates, sulfonamides).
On irritated or broken mucous membranes.
When Using: Avoid eyes/nostrils; rinse if contact occurs.
Do not swallow immediately; allow throat numbing.
Avoid eating/drinking for 1 hour post-use to prevent choking.
Seek immediate medical help for excessive numbness, breathing issues, dizziness, or bluish skin.

Stop Use and Consult a Doctor If: Severe burning, swelling, irritation, rash, or prolonged numbing (>1-2 hours) occurs.

Ask a Doctor Before Use If: You have respiratory, heart, or liver issues, are pregnant/breastfeeding, or take interacting medications.

Keep Out of Reach of Children: Contact Poison Control

(1-800-222-1222) if swallowed.

• Other Warnings: Overuse may increase methemoglobinemia risk.
Avoid continuous/large

Keep out of reach of childeren.Contact Poison Control if

Directions

Adults (18+): 1-2 sprays to throat, max 3-4 per use.

Allow 30-60 seconds before swallowing; wait 1-2 minutes for full effect. Numbing fades within 1 hour. Max 4 sprays/session, 12 sprays/24 hours.

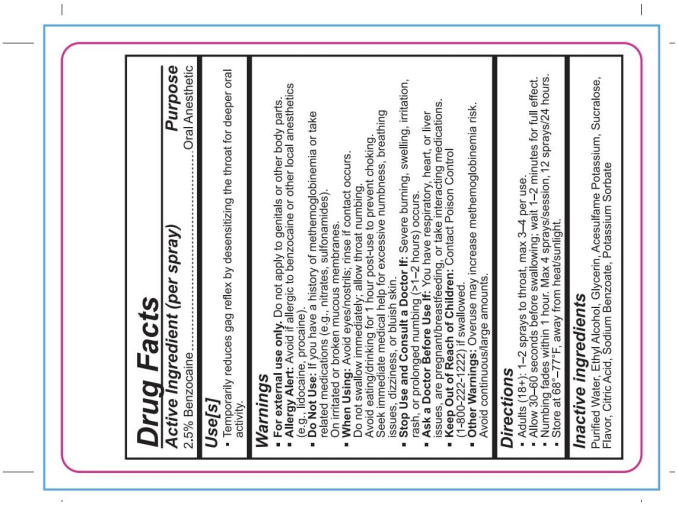
Store at 68°-77°F, away from heat/sunlight.

Other information

Inactive ingredients

Purified Water, Ethyl Alcohol, Glycerin, Acesulfame Potassium, Sucralose, Flavor, Citric Acid, Sodium Benzoate, Potassium

Package Label - Principal Display Panel



BVIBE SPEARMINT DEEP THROAT

benzocaine numbing spray liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77632-161
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	25 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ACESULFAME POTASSIUM (UNII: 23OV73Q5G9)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

CITRIC ACID (UNII: 2968PHW8QP)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SPEARMINT (UNII: J7I2T6IV1N)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77632-161-11	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/15/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	07/15/2025	

Labeler - Private Label Productions LLC (046278265)

Revised: 8/2025 Private Label Productions LLC