

**MOUTH RINSE- eucalyptol, menthol, methylsalicylate, thymol mouthwash  
Kroger**

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**Spring Mint Antiseptic Mouthrinse  
072AL rev2-072AN**

**Active ingredients**

Eucalyptol 0.092%

Menthol 0.042%

Methyl salicylate 0.060%

Thymol 0.064%

**Purpose**

Antigingivitis, Antiplaque

**Use**

helps control plaque that leads to gingivitis

**Warnings**

For this product

**Do not use**

If you have painful or swollen gums, pus from the gum line, loose teeth or increased spacing between the teeth. See your dentist immediately. These may be signs of periodontitis, a serious form of gum disease.

**Stop use and ask a dentist if**

gingivitis, bleeding, or redness persists for more than 2 weeks

**Keep out of reach of children**

If more than used for rinsing is accidentally swallowed, get medical help or contact a Poison Control Center right away.

**Directions**

Adults and children 12 years of age and older - vigorously swish 20 mL (2/3 FL OZ or 4 teaspoonfuls) between teeth for 30 seconds then spit out; do not swallow

children under 12 years of age - consult a dentist or doctor

- this rinse is not intended to replace brushing or flossing

### **Other information**

Cold weather may cloud this product. Its antiseptic properties are not affected. Store at room temperature (59°- 77°F).

### **Inactive ingredients**

water, alcohol 21.6%, sorbitol, poloxamer 407, benzoic acid, sodium saccharin, flavor, sodium citrate, yellow 10, green 3

### **Questions or comments?**

1-800-632-6900

### **Disclaimer**

\*Listerine is a registered trademark of Johnson & Johnson Healthcare Products, Skillman, NJ 08558. Johnson & Johnson Healthcare Products is not affiliated with The Kroger Co. or this product.

### **Adverse Reactions**

DISTRIBUTED BY THE KROGER CO.

CINCINNATI, OHIO 45202

QUALITY GUARANTEE

[www.kroger.com](http://www.kroger.com)

DSP-TN-21091

DSP-NO-20087

### **Principal display panel**

COMPARE to the active ingredients of FRESH BURST® LISTERINE®

\*See Back Label

Kroger

Mint Burst

ANTISEPTIC

MOUTH RINSE

ANTGINGIVITIS/ANTIPLAQUE

- Kills Germs That Cause Bad Breath, Plaque & the Gum Disease Gingivitis
- For Fresher Breath

ADA Accepted

American Dental Association

- Helps prevent and reduce plaque
- Helps prevent and reduce gingivitis

SEALED WITH PRINTED NECKBAND FOR YOUR PROTECTION

16.9 FL OZ (500 mL)

COMPARE to the active ingredients of FRESH BURST® LISTERINE®

\*See Back Label



Mint Burst

# ANTISEPTIC MOUTH RINSE

## ANTIGINGIVITIS/ANTIPLAQUE

- Kills Germs That Cause Bad Breath, Plaque & the Gum Disease Gingivitis
- For Fresher Breath



SEALED WITH PRINTED NECKBAND FOR YOUR PROTECTION



16.9 FL OZ (500 mL)

L0002130FF

## MOUTH RINSE

eucalyptol, menthol, methylsalicylate, thymol mouthwash

### Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:30142-072

**Route of Administration** ORAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>EUCALYPTOL</b> (UNII: RV6J6604TK) (EUCALYPTOL - UNII:RV6J6604TK)	EUCALYPTOL	0.92 mg in 1 mL
<b>MENTHOL</b> (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.42 mg in 1 mL
<b>METHYL SALICYLATE</b> (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	0.60 mg in 1 mL
<b>THYMOL</b> (UNII: 3J50XA376E) (THYMOL - UNII:3J50XA376E)	THYMOL	0.64 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>POLOXAMER 407</b> (UNII: TUF2IVW3M2)	
<b>BENZOIC ACID</b> (UNII: 8SKN0B0MIM)	
<b>SACCHARIN SODIUM</b> (UNII: SB8ZUX40TY)	
<b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)	
<b>D&amp;C YELLOW NO. 10</b> (UNII: 35SW5USQ3G)	
<b>FD&amp;C GREEN NO. 3</b> (UNII: 3P3ONR6O1S)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30142-072-77	500 mL in 1 BOTTLE, PLASTIC; Type 1: Convenience Kit of Co-Package	11/11/1989	
2	NDC:30142-072-86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/11/1989	
3	NDC:30142-072-12	1500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/11/1989	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	11/11/1989	

**Labeler** - Kroger (006999528)

**Registrant** - Vi Jon, LLC (790752542)

## Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		790752542	manufacture(30142-072)

## Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		088520668	manufacture(30142-072)

Revised: 10/2023

Kroger