ANTISEPTIC MOUTH RINSE- eucalyptol mouthwash
Walgreens Co.

Spring Mint Antiseptic Mouthrinse
072.003/072AL rev 2-072AN

TEP

SEALED WITH PRINTED NECKBAND FOR YOUR PROTECTION

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. It antiseptic properties are not affected. Store at room temperature (59º - 77ºF)

Inactive ingredients

water, alcohol 21.6%, sorbitol solution, flavor, poloxamer 407, benzoic acid, sodium saccharin, sodium citrate, yellow 10, green3

Disclaimer

Our pharmacists recommend the Walgreens brand. We invite you to compare to nation brands.

*This product is not manufactured or distributed by Johnson & Johnson, owner of the registered tradearks Listerine and Freshburst.

DISTRIBUTED BY: WALGREEN CO.

200 WILMOT RD., DEERFIELD, IL 60015

100% SATISFACTION GUARANTEED

walgreens.com

MADE IN THE U.S.A. WITH U.S. AND FOREIGN COMPONENTS

principal display panel

WALGREENS - PHARMACIST RECOMMENDED

Walgreens

Compare to the active ingredients in Listine Freshburts*

Mouth Rinse

ANTIGINGIVITIS/ANTIPLAQUE

Antiseptic

- Kills germs that cause bad breath, plaque & the gum disease gingivitis

ADA Accepted

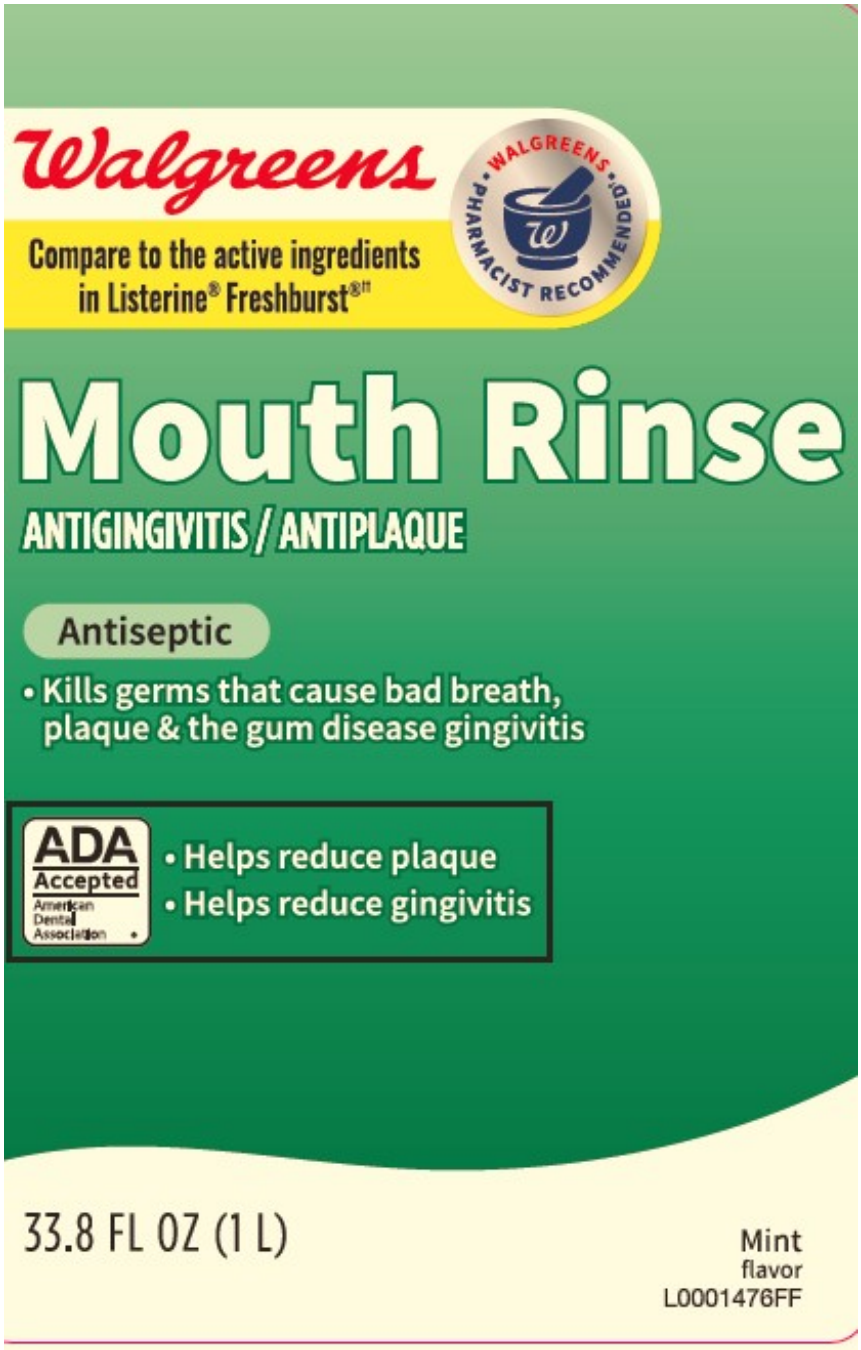
American

Dental Association

- Helps reduce plaque
- Helps reduce gingivitis

33.8 FL OZ (1 L)

Mint flavor



ANTISEPTIC MOUTH RINSE eucalyptol mouthwash

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-0072
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
EUCALYPTOL (UNII: RV6J6604TK) (EUCALYPTOL - UNII:RV6J6604TK)	EUCALYPTOL	0.92 mg in 1 mL
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.42 mg in 1 mL
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	0.6 mg in 1 mL
THYMOL (UNII: 3J50XA376E) (THYMOL - UNII:3J50XA376E)	THYMOL	0.64 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	MINT (Spring Mint)	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0072-77	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2007	
2	NDC:0363-0072-86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2007	
3	NDC:0363-0072-12	1500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2007	
4	NDC:0363-0072-13	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2007	
5	NDC:0363-0072-88	2000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2007	

6	NDC:0363-0072-69	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2007	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/30/2007	

Labeler - Walgreens Co. (008965063)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(0363-0072)

Revised: 7/2024 Walgreens Co.