ANTISEPTIC MOUTH RINSE- eucalyptol, menthol, methyl salicylate, thymol mouthwash CDMA

Quality Choice 318.003/318AH-AI Antiseptic Mouth Rinse

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

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

- do not swallow
- 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 26.9%, benzoic acid, poloxamer 407, sodium benzoate, caramel

Disclaimer

*This product is not manufactured or distributed by Johnson & Johnson Healthcare Products, distributor of Original Listerine Antiseptic Mouthwash

Adverse reaction

Distributed by C.DM.A., Inc

43157 W 9 Mile Rd

Novi, MI 48375

www.qualitychoice.com

Questions: 800-935-2362

100% OC SATISFACTION GUARANTEED

Principal display panel

Sealed With Printed Neckband For Your Protection

QC

QUALITY

CHOICE

Compare to Original LISTERINE*

Antiseptic

Mouth Rinse

Antigingivitis/Antiplaque

Kills Germs That Cause:

Bad Breath

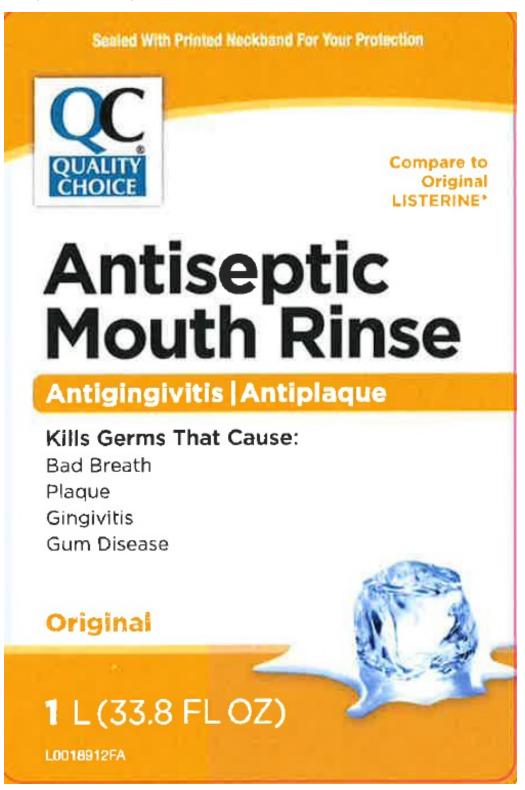
Plaque

Gingivitis

Gum Disease

Original

1 L (33.8 FL OZ)



eucalyptol, menthol, methyl salicylate, thymol mouthwash

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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)		
BENZOIC ACID (UNII: 85KN0B0MIM)		
POLOXAMER 407 (UNII: TUF2IVW3M2)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
CARAMEL (UNII: T9D99G2B1R)		

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:83324- 180-33	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/02/2024	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	505G(a)(3)	07/02/2024		

Labeler - CDMA (011920774)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Na me	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(83324-180)

Revised: 7/2024 CDMA