

**ANTISEPTIC MOUTHRINSE- eucalyptol, menthol, methyl salicylate,
thymol mouthwash**
Chain Drug Marketing Association

Quality Choice 072.003/072AN
Spring Mint Antiseptic Mouthrinse

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 twice a day 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, D&C yellow no. 10, FD&C green no.3

Disclaimer

*This product is not manufactured or distributed by Kenvue, Inc., distributor of FreshBurst® Listerine Antiseptic Mouthwash.

Adverse Reactions

QC 100% SATISFACTION GUARANTEED®

Distributed by CDMA, Inc.

Novi, MI 48375

www.qualitychoice.com

Questions: 800-935-2362

DSP-TN-21091

DSP-MO-20087

Principal display panel

Sealed With Printed Neckband For Your Protection

NDC 83324-181-33

QC®

QUALITY

CHOICE

Compare to FreshBurst® Listerine®*

Antiseptic

Mouth Rinse

Antigingivitis/Antiplaque

Kills Germs That Cause:

Bad Breath

Plaque

Gingivitis

Gum Disease

Spring Mint[®]

1 L (1.05 QT) 33.8 FL OZ

Sealed With Printed Neckband For Your Protection

NDC 83324-781-33



Compare to
FreshBurst[®]
LISTERINE^{®*}

Antiseptic Mouth Rinse

Antigingivitis | Antiplaque

Kills Germs That Cause:
Bad Breath
Plaque
Gingivitis
Gum Disease



Spring Mint[®]

**1 L (1.05 QT)
33.8 FL OZ**

L0018913FB

Drug Facts

Active ingredients

Eucalyptol 0.092%, Menthol 0.042% Antigingivitis, Antiplaque
Methyl salicylate 0.060%, Thymol 0.064% Antigingivitis, Antiplaque

Purpose

Use helps control plaque that leads to gingivitis

Warnings

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 and older	vigorously swish 20 mL (2/3 FL OZ or 4 teaspoonfuls) between teeth twice a day 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

*This product is not manufactured or distributed by Kenvue, Inc., distributor of FreshBurst® Listerine® Antiseptic Mouthwash.



Distributed by CDMA, Inc.
Novi, MI 48375
www.qualitychoice.com
Questions: 800-935-2362



DSP-142 091
DSP-MC-20087

L0018913BB

ANTISEPTIC MOUTHRINSE

eucalyptol, menthol, methyl salicylate, thymol mouthwash

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83324-181
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 (UNII: 506T60A25R)	
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-181-33	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 - Chain Drug Marketing Association (011920774)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091514	manufacture(83324-181)