ANTISEPTIC- eucalyptol, menthol, methyl salicylate, thymol mouthwash
Vi Jon, LLC

Swan 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 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 ingredents

water, alcohol 21.6%, sorbitol solution, flavor, poloxamer 407, benzoic acid, sodium saccharin, sodium citrate, D&C yellow no. 10, FD&C green no. 3

Disclaimer

This product is not manufactured or distributed by Johnson & Johnson Healthcare Products, distributor Listerine ®.

Adverse reaction

Distributed By:

Vi-Jon, LLC

One Swan Drive

Smyrna, TN 37167

Principal display panel

Sealed With Printed Neckband For Your Protection

swan®

ANTISEPTIC

Mouth Rinse

spring mint®

Kills Germs that Cause Bad Breath, Plaque & the Gum Disease Gingivits.

Compare to the active ingredients in FreshBurst® Listerine®*

ADA ACCEPTED

AMERICAN DENTAL ASSOCIATION
- Helps reduce plaque
- Helps reduce gingivitis

1 LITER (33.8 FL OZ)

ANTISEPTIC

eucalyptol, menthol, methyl salicylate, thymol mouthwash

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0869-0072
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 (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:0869-0072-21	89 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/1988	03/13/2024
2	NDC:0869-0072-88	2000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/1988	03/13/2024
3	NDC:0869-0072-69	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/1988	
4	NDC:0869-0072-77	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/1988	
5	NDC:0869-0072-86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/1988	
6	NDC:0869-0072-19	94 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/1988	03/31/2017
7	NDC:0869-0072-50	710 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/1988	03/13/2024
8	NDC:0869-0072-13	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/1988	03/13/2024
9	NDC:0869-0072-12	1500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/1988	06/06/2021

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	07/20/1988	

Labeler - Vi Jon, LLC (088520668)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(0869-0072)

Revised: 3/2024 Vi Jon, LLC