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

Swan 664.003/664AT rev2-AU
Blue 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 (59º-77ºF).

Inactive ingredients

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

ADA Council Statement

The ADA Council on scientific affairs Acceptance of Swan Blue Mint Antiseptic mouth rinse is based on its finding that the product is effective in helping to prevent and reduce gingivitis and plque above the gumline, when used as directed

Disclaimer

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

Distributed by: Vi-Jon

One Swan Drive

Smyrna, TN 37167

DSP-TN-15000

DSP-MO-34 SDS-TN-15012

principal display panel

Sealed With Printed Neckband For Your Protection

NDC 0869-0664-77

Swan ®

ANTISEPTIC

MOUTH RINSE

ice mint ®

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

Compare to active ingredients of Listerine ®*

ADA Accepted

American Dental Association

500 mL (16.9 FL OZ)

ANTISPETIC

eucalyptol, menthol, methyl salicylate, thymol mouthwash

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0869-0664
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 BENZOATE (UNII: OJ245FE5EU)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0869-0664-86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1992	05/12/2025
2	NDC:0869-0664-77	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1992	
3	NDC:0869-0664-69	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1992	05/12/2025
4	NDC:0869-0664-88	2000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1992	04/25/2024
5	NDC:0869-0664-13	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1992	09/27/2015
6	NDC:0869-0664-12	1500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1992	
7	NDC:0869-0664-19	94 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1992	12/01/2019

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	07/15/1992	

Labeler - Vi-Jon, LLC (088520668)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(0869-0664)

Revised: 5/2025 Vi-Jon, LLC