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

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Swan Mountain Falls 072.003/072AN

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

ADA council statement

“The ADA Council on Scientific Affairs’ Acceptance of Swan Spring Mint® Antiseptic Mouth Rinse is based on its finding that the product is effective in helping to prevent and reduce gingivitis and plaque above the gumline, when used as directed.”

safety information

SEALED WITH PRINTED NECKBAND FOR YOUR PROTECTION

disclaimer

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

DSP-TN-1500 DSP-MO-34 SDS-TN-15012

Adverse Reactions

Distributed By: Vi-Jon, LLC

8515 Page Ave., St. Louis, MO 63114

Questions or Comments? 1-888-593-0593

principal display panel

NDC 0869-0072-69

Sealed With Printed Neckband For Your Protection

SWAN

ANTISEPTIC

MOUTH RINSE

spring mint

Kills Germs that Cause

Bad Breath, Plaque & the

Gum Disease Gingivitis

Compare to FreshBurst Listerin

ADA

Accepted

American

Dental

Association

250 mL (8.5 FL OZ)

principal display panel

Mountain

Falls

Compare to Listerine

improves oral hygiene

for daily mouth care

kills germs that cause bad breath, plaque and gingivitis gum disease

freshens breath

antiseptic

mouth rinse

antigingivitis/antiplaque

spring mint

500 mL (16.9 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	0.60 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	
2	NDC:0869-0072-88	2000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/1988	
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	
7	NDC:0869-0072-50	710 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/1988	
8	NDC:0869-0072-13	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/1988	
9	NDC:0869-0072-12	1500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/1988	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not	part356	07/20/1988	

final	part356	07/20/1988	

Labeler - Vi Jon, LLC (790752542)

Registrant - Vi Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		790752542	manufacture(0869-0072)

Revised: 2/2022 Vi Jon, LLC