ANTISEPTIC MOUTHRINSE- eucalyptol, menthol, methyl salicylate, thymol mouthwash
Wal-Mart Stores, Inc.

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.

Spring Mint Antiseptic Mouthrinse
072.003/072AL rev 2

TEP

Sealed with printed neckband for your protection

Active ingredients

Eucalyptol 0.092%, Menthol 0.042%, Methyl sailcylate 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 incresed 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 rednss 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 under 12 years if 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 propeties are not affected. Store at room temperature (59° -77° F)

Inactive ingredients

water, alcohol 21.6 %, sorbitol solution, flavor, poloxamer 407, benzoic acid, sodium saccharin, sodium citrate, D&C yellow 10, FD&Cgreen 3

Satisfaction guaranteed

- Or we'll replace it or give you your money back. For questions or comments or to report an undesired reaction or side effect, please call 1-888-287-1915.

Adverse Reactions Section

DISTRIBUTED BY: Wal-Mart Stores, Inc., Bentonville, AR 72716

*This product is not manufactured or distributed by Johnson & Johnson Healthcare Products, owner of the registered mark FreshBurst Listerine.

how2recycle.info

Discard Seal, Empty & Replace Cap

Plastic Bottle

Principal Display Panel

NDC: 49035-072-12

equate

Compare to FreshBurst Listerine Active Ingredients*

Antiseptic Mouthrinse

Kills Germs that Cause:
- Plaque
- Gingivitis
- Bad breath

ADA Accepted

American Dental Association
- Helps reduce plaque
- Helps reduce gingivitis

Spring Mint

Made in the USA

with 90% OR MORE US PARTS

Factory Certified

1.5 LITERS (50.7 FL OZ)

ANTISEPTIC MOUTHRINSE

eucalyptol, menthol, methyl salicylate, thymol mouthwash

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49035-072
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

MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	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:49035-072-69	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/11/1989	
2	NDC:49035-072-77	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/11/1989	
3	NDC:49035-072-86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/11/1989	
4	NDC:49035-072-12	1500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/11/1989	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	11/11/1989	

Labeler - Wal-Mart Stores, Inc. (051957769)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(49035-072)

Establishment

Name	Address	ID/FEI	Business Operations

Vi-Jon, LLC		088520668	manufacture(49035-072)