FAMILY WELLNESS ORIGINAL AMBER- eucalyptol, menthol, methyl salicylate, thymol mouthwash

Family Dollar Services 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.

Family Wellness Original Amber Mouthwash

Drug Facts

Active ingredients

Eucalyptol 0.092%, Menthol 0.042%

Methyl Salicylate 0.060%, Thymol 0.064%

Purpose

Antigingivitis, Antiplaque

Antigingivitis, Antiplaque

*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 of age and older: vigorously swish 20 mL (2/3 FL OS 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 $(59^{\circ} - 77^{\circ}F)$

Inactive ingredients water, alcohol 21.6%, benzoic acid, poloxamer 407, sodium benzoate, caramel.

DISTRIBUTED BY: Midwood Brands, LLC

500 Volvo Parkway, Chesapeake, VA 23320

Made in Canada

* This product is not manufactured or distributed by Johnson & Johnson, distributor of Listerine®.

COMPARE TO LISTERINE® ORIGINAL*

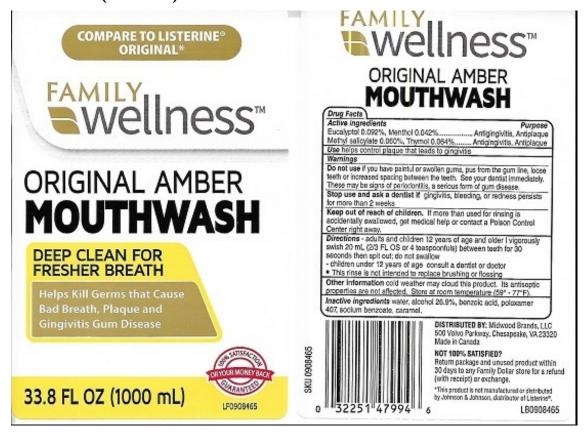
FAMILY wellnessTM

ORIGINAL AMBER

MOUTHWASH

DEEP CLEAN FOR FRESHER BREATH

Helps Kill Germs that Cause Bad Breath, Plaque and Gingivitis Gum Disease 33.8 FL OZ (1000 mL)



FAMILY WELLNESS ORIGINAL AMBER

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Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55319-232
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
EUCALYPTOL (UNII: RV6J6604TK) (EUCALYPTOL - UNII:RV6J6604TK)	EUCALYPTOL	0.092 g in 100 mL
MENTHOL (UNII: L7T10 EIP3A) (MENTHOL - UNII:L7T10 EIP3A)	MENTHOL	0.042 g in 100 mL
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	0.06 g in 100 mL
THYMOL (UNII: 3J50XA376E) (THYMOL - UNII:3J50XA376E)	THYMOL	0.064 g in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
SODIUM BENZOATE (UNII: OJ245FE5EU)	
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
BENZOIC ACID (UNII: 8 SKN0 B0 MIM)	

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:55319-232- 86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/20/2019		
2	NDC:55319-232- 23	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/20/2019		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part356	06/20/2019		

Labeler - Family Dollar Services Inc. (024472631)

Revised: 1/2020 Family Dollar Services Inc.