

**ANTISEPTIC- eucalyptol, menthol, methyl salicylate, thymol liquid
WALMART INC. (see also Equate)**

Kalaya Breath Refresh Antiseptic Oral Rinse-updated label for NDC

Active ingredient

Eucalyptol (0.092%)

Menthol (0.042%)

Methyl Salicylate (0.060%)

Thymol (0.064%)

Purposes

Antiplaque/antigingivitis

Uses

helps prevent and reduce

- plaque
- gingivitis

Warning

Do not use

in children under 12 years of age.

Stop use and ask a dentist

- gingivitis, bleeding, or redness persists for more than 2 weeks
- you have painful or swollen gums, pus from the gum line, loose teeth, or increasing spacing between the teeth. These may be signs or symptoms of periodontitis, a serious form of gum disease.

Keep out of reach of children under 6 years of age

If more than used for rinsing is accidentally swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away

Other information

- store at room temperature
- cold weather may cloud this product. Its antiseptic properties are not affected.
- this rinse is not intended to replace brushing or flossing

Directions

- adults and children 12 years of age and older: vigorously swish 20 milliliters of rinse between your teeth twice a day for 30 seconds and then spit out. Do not swallow the rinse.
- children 6 years to under 12 years of age: supervise use
- children under 6 years of age: do not use

Inactive ingredients

water, alcohol (21.6% v/v), sorbitol, poloxamer 407, flavor, sodium benzoate, benzoic acid, sodium saccharin, green #3

Questions or comments

1-888-287-1915

Principal display panel

Compare to Listerine® Cool Mint® active ingredient

Antiseptic Mouthwash

Eucalyptol (0.092%)

Menthol (0.042%)

Methyl salicylate (0.060%)

Thymol (0.064%)

Antiplaque/antigingivitis

- Kills germs that cause plaque, gingivitis, bad breath
- Help to control and prevent plaque that leads to gingivitis

Blue Mint

L (QT PT FL OZ)

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

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL AROUND OR UNDER CAP IS BROKEN OR MISSING.

DISTRIBUTED BY: Walmart Inc.,

Bentonville, AR 72716

Package label

NDC 79903-151-75

Antiseptic Mouthwash

Eucalyptol (0.092%)
Menthol (0.042%)
Methyl salicylate (0.060%)
Thymol (0.064%)

Antiplaque/antigingivitis

- Kills germs that cause plaque, gingivitis, bad breath
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Blue Mint

1.5 L (1 QT 1 PT 2.7 FL OZ)

PLD-A755A LB008986

Drug Facts

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Uses helps prevent and reduce ■ plaque ■ gingivitis

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Questions or comments? 1-888-287-1915

Satisfaction guaranteed - Or we'll replace it or give you your money back. For questions or comments please call 1-888-287-1915.

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL AROUND BOTTLE OR UNDER CAP IS BROKEN OR MISSING.



DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716
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PLD-A755A LB008986

EQUATE Antiseptic Mouthwash

ANTISEPTIC

eucalyptol, menthol, methyl salicylate, thymol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79903-151
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)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	

Product Characteristics

Color	blue	Score	
Shape		Size	
Flavor	MINT	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-151-75	1.5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/30/2022	
2	NDC:79903-151-50	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/30/2022	
3	NDC:79903-151-99	2 in 1 PACKAGE	12/30/2022	
3		1.5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M022	12/30/2022	

Labeler - WALMART INC. (see also Equate) (051957769)

Revised: 6/2024

WALMART INC. (see also Equate)