

LISTERINE COOL MINT ANTISEPTIC- eucalyptol, menthol, methyl salicylate, thymol mouthwash
Kenvue Brands LLC

Listerine Cool Mint Antiseptic Mouthwash

Drug Facts

Active ingredient

Eucalyptol 0.092%

Menthol 0.042%

Methyl Salicylate 0.060%

Thymol 0.064%

Purpose

Antiplaque/antigingivitis

Uses

helps prevent and reduce: • plaque • gingivitis

Warnings

Do not use in children under 12 years of age

Ask a dentist

Ask a dentist if symptoms persist, new symptoms appear, or conditions worsen after regular use

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

- rinse full strength for 30 seconds with 20 mL (2/3 fluid ounce or 4 teaspoonfuls) morning and night
- do not swallow

Other information

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

Inactive ingredients

Water, Alcohol (21.6%), Sorbitol, Poloxamer 407, Benzoic Acid, Sodium Saccharin, Sodium Benzoate, Flavor, Green 3

Questions?

call toll-free **888-222-0182** or **215-273-8755** (collect)

Distributed by:

Kenvue Brands LLC

Summit, NJ 07901

PRINCIPAL DISPLAY PANEL - 1 L Bottle Label

ANTIGINGIVITIS/ANTIPLAQUE MOUTHWASH

LISTERINE®

COOL MINT®

ANTISEPTIC

FOR A FRESHER &

CLEANER MOUTH THAN

BRUSHING ALONE

ADA

Accepted

American

Dental

Association®

1.0 L (1 Qt 1.8 Fl Oz)



Drug Facts

| | |
|---|---|
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|---|---|

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Do not use if printed band around cap is broken or missing.

ADA Accepted
American Dental Association

- Helps prevent and reduce plaque
- Helps prevent and reduce gingivitis

Distributed by:
Kenvue Brands LLC
Summit, NJ 07901
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Pat. www.kenvuepats.com

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PLASTIC BOTTLE

Empty & Replace Cap

how2recycle.info

EXP LOT

LISTERINE COOL MINT ANTISEPTIC

eucalyptol, menthol, methyl salicylate, thymol mouthwash

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|----------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:69968-0791 |
| Route of Administration | ORAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|-----------------|
| 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 |
| 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 |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| WATER (UNII: 059QF0KO0R) | |
| ALCOHOL (UNII: 3K9958V90M) | |
| POLOXAMER 407 (UNII: TUF2IVW3M2) | |
| BENZOIC ACID (UNII: 8SKN0B0MIM) | |

| | |
|--|--|
| SORBITOL (UNII: 506T60A25R) | |
| SODIUM BENZOATE (UNII: OJ245FE5EU) | |
| SACCHARIN SODIUM (UNII: SB8ZUX40TY) | |
| FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:69968-0791-1 | 1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 03/21/2023 | |
| 2 | NDC:69968-0791-9 | 95 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 03/21/2023 | |
| 3 | NDC:69968-0791-3 | 250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 03/21/2023 | |
| 4 | NDC:69968-0791-2 | 1500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 03/21/2023 | |
| 5 | NDC:69968-0791-4 | 3700 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 03/21/2023 | |
| 6 | NDC:69968-0791-5 | 500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 03/21/2023 | |
| 7 | NDC:69968-0791-6 | 2 in 1 PACKAGE | 03/21/2023 | |
| 7 | | 1500 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 | 03/21/2023 | |

Labeler - Kenvue Brands LLC (118772437)

Revised: 3/2025

Kenvue Brands LLC