

**LISTERINE COOL MINT ANTISEPTIC- eucalyptol, menthol, unspecified form, methyl salicylate, and thymol liquid
Johnson & Johnson Consumer 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.

Listerine Cool Mint Antiseptic

Drug Facts

Active ingredients

Eucalyptol 0.092%

Menthol 0.042%

Methyl Salicylate 0.060%

Thymol 0.064%

Purposes

Antiplaque/antigingivitis

Uses

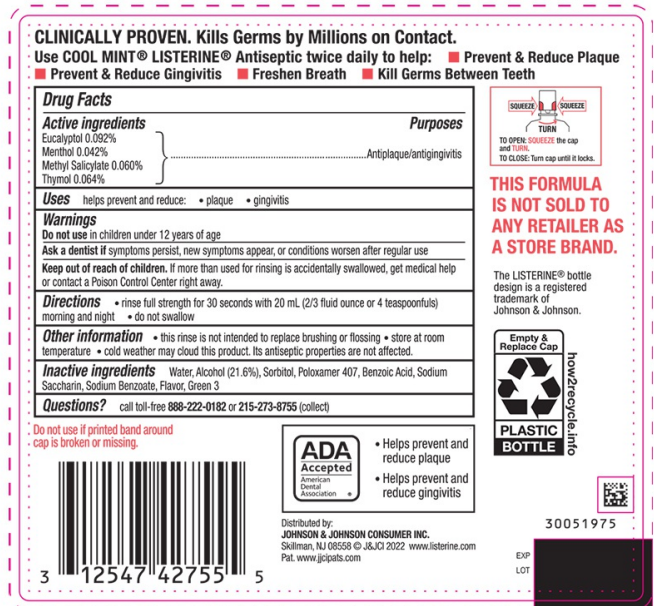
Helps prevent and reduce:

- Plaque
- Gingivitis

Warnings

**KILLS 99.9% OF GERMS
THAT CAUSE BAD BREATH,
PLAQUE & GINGIVITIS**

30040767



Do not use in children under 12 years of age

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 or Comments?

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

Distributed by:

JOHNSON & JOHNSON CONSUMER INC.

Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 1.5 L Bottle Label ANTISEPTIC

LISTERINE®

COOL MINT®

FOR A FRESHER &

CLEANER MOUTH THAN

BRUSHING ALONE

ADA

Accepted

American

Dental

Association®

1.5L (1 Qt 1 Pt 2.7 fl oz)

**KILLS 99.9% OF GERMS
THAT CAUSE BAD BREATH,
PLAQUE & GINGIVITIS**

30040767



CLINICALLY PROVEN. Kills Germs by Millions on Contact.
Use **COOL MINT® LISTERINE®** Antiseptic twice daily to help: ■ Prevent & Reduce Plaque ■ Prevent & Reduce Gingivitis ■ Freshen Breath ■ Kill Germs Between Teeth

Drug Facts	
Active ingredients	Purposes
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Menthol 0.042%	
Methyl Salicylate 0.060%	
Thymol 0.064%	
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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:
JOHNSON & JOHNSON CONSUMER INC.
Skillman, NJ 08553 © J&J 2022 www.listerine.com Pat. www.jjcpats.com

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30051975

EXP
LOT

LISTERINE COOL MINT ANTISEPTIC

eucalyptol, menthol, unspecified form, methyl salicylate, and thymol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0550
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, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	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
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WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
SORBITOL (UNII: 506T60A25R)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0550-9	95 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/03/2018	
2	NDC:69968-0550-2	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/03/2018	
3	NDC:69968-0550-5	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/03/2018	
4	NDC:69968-0550-1	946.353 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/03/2018	
5	NDC:69968-0550-4	1419.529 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/03/2018	
6	NDC:69968-0550-3	3700 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2019	
7	NDC:69968-0550-8	2 in 1 PACKAGE	11/24/2020	
7	NDC:69968-0550-4	1419.529 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part356	12/03/2018	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 1/2023

Johnson & Johnson Consumer Inc.