LISTERINE CLINICAL SOLUTIONS GUM HEALTH ICY MINT- eucalyptol, menthol, methyl salicylate, thymol mouthwash
Johnson & Johnson Consumer Inc.

Listerine Clinical Solutions Gum Health Icy Mint

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

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, Sorbitol, Poloxamer 407, Flavor, Zinc Chloride, Benzoic Acid, Sodium Benzoate, Sodium Saccharin, Sucralose, Red 40

Questions?

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

Distributed by:

JOHNSON & JOHNSON CONSUMER INC.

Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 1 L Bottle Label

ANTIGINGIVITIS / ANTIPLAQUE MOUTHWASH

LISTERINE ®

CLINICAL

SOLUTIONS

ANTISEPTIC

GUM HEALTH

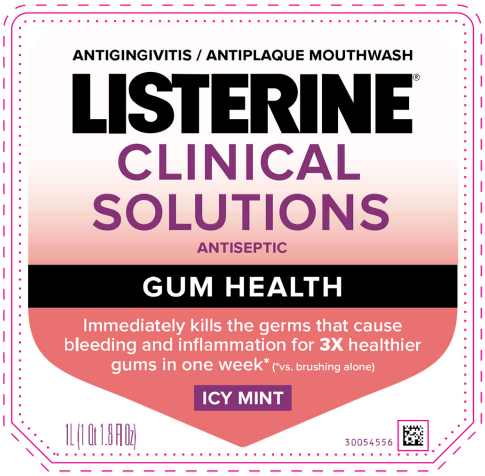
Immediately kills the germs that cause

bleeding and inflammation for **3X** healthier

gums in one week* (*vs. brushing alone)

ICY MINT

1L (1 Qt 1.8 Fl Oz)



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eucalyptol, menthol, methyl salicylate, thymol mouthwash

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0818
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)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
ZINC CHLORIDE (UNII: 86Q357L16B)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0818-9	95 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/16/2023	
2	NDC:69968-0818-5	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/16/2023	
3	NDC:69968-0818-1	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/16/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M022	10/16/2023	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 10/2023 Johnson & Johnson Consumer Inc.