

LISTERINE GUM THERAPY GLACIER MINT- eucalyptol, menthol, methyl salicylate, thymol mouthwash
Johnson & Johnson Consumer Inc.

Listerine Gum Therapy Mouthwash

Glacier Mint

Drug Facts

Active ingredient	Purpose
Eucalyptol (0.092%)	Antiplaque/antigingivitis
Menthol (0.042%)	Antiplaque/antigingivitis
Methyl Salicylate (0.060%)	Antiplaque/antigingivitis
Thymol (0.064%)	Antiplaque/antigingivitis

Use

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 (21.6% v/v), Sorbitol, Poloxamer 407, Benzoic Acid, Zinc Chloride,

Sodium Benzoate, Sucralose, Flavor, Sodium Saccharin, Blue 1

Questions?

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

Distributed by:

JOHNSON & JOHNSON CONSUMER INC.

Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 1.0 L Bottle Label

ANTINGIVITIS / ANTIPLAQUE MOUTHWASH

LISTERINE®

GUM THERAPY

4X

HEALTHIER*

**HELPS REVERSE SIGNS OF
EARLY GUM DISEASE: REDNESS,
BLEEDING & INFLAMMATION**

GLACIER MINT

1.0 L (1 Qt 1.8 Fl Oz)

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1.0 L (1 Qt 1.8 Fl Oz) **GLACIER MINT**

30053782



CLINICALLY PROVEN. Helps reverse signs of early gum disease in 3 weeks.
LISTERINE® Gum Therapy cleans below the gumline where brushing can't reach to kill harmful bacteria that cause red, inflamed, and bleeding gums.

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..... Antiplaque/antigingivitis

Uses helps prevent and reduce: • plaque • gingivitis

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THIS FORMULA IS NOT SOLD TO ANY RETAILER AS A STORE BRAND.

The LISTERINE® bottle design is a registered trademark of Johnson & Johnson.



30053753



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



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

Distributed by:
JOHNSON & JOHNSON CONSUMER INC. EXP
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Pat. www.jjci-pats.com

LISTERINE GUM THERAPY GLACIER MINT			
eucalyptol, menthol, methyl salicylate, thymol mouthwash			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0790
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		

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 BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0790-5	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/13/2023	
2	NDC:69968-0790-1	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/13/2023	
3	NDC:69968-0790-9	95 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/13/2023	
4	NDC:69968-0790-2	1500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/01/2023	
5	NDC:69968-0790-3	2 in 1 TRAY	03/01/2024	
5		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	02/13/2023	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 7/2024

Johnson & Johnson Consumer Inc.