

**LISTERINE GUM THERAPY ANTISEPTIC- eucalyptol, menthol, unspecified form, methyl salicylate, and thymol mouthwash
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 GUM THERAPY ANTISEPTIC

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

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 20mL (2/3 fluid ounce or 4 teaspoonfuls) morning and night
- do not swallow

Other information

- 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

**ANTISEPTIC
LISTERINE
GUM THERAPY
4X
HEALTHIER***

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

GLACIER MINT
1.0 L (1 Qt 1.8 Fl Oz)



**FOR 4X HEALTHIER
GUMS IN 3 WEEKS***

***VS BRUSHING ALONE**

30045649



ANTISEPTIC

LISTERINE® **GUM THERAPY**



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

30045648

1.0 L (1 Qt 1.8 Fl Oz)

GLACIER MINT



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.

Drug Facts

Active ingredients

Purposes



Active Ingredients

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- Menthol (0.042%)
- Methyl Salicylate (0.060%)
- Thymol (0.064%)

Antiplaque/antigingivitis

Uses helps prevent and reduce: • plaque • gingivitis

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TURN
TO OPEN: SQUEEZE the cap and **TURN**,
TO CLOSE: Turn cap until it locks.

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

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



30045647

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



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JOHNSON & JOHNSON CONSUMER INC.
Skillman, NJ 08558
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www.listerine.com

EXP
LOT

LISTERINE GUM THERAPY ANTISEPTIC

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

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0604
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: TUF21VW3M2)	
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:69968-0604-5	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/06/2020	09/26/2023
2	NDC:69968-0604-1	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/06/2020	09/27/2023
3	NDC:69968-0604-9	95 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/06/2020	09/26/2023

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part356	01/06/2020	09/27/2023

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 3/2023

Johnson & Johnson Consumer Inc.