LISTERINE ULTRACLEAN ANTISEPTIC COOL MINT- 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 Ultraclean Antiseptic
Cool 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

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

- store at room temperature
- cold weather may cloud this product. Its antiseptic properties are not affected.

Inactive ingredients

water, alcohol (21.6%), sorbitol, flavor, poloxamer 407, benzoic acid, zinc chloride, sodium benzoate, sucralose, sodium saccharin, green 3

Questions?

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

Distributed by:
JOHNSON & JOHNSON CONSUMER INC.
Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 500 mL Bottle Label

ANTISEPTIC
LISTERINE®
ULTRACLEAN

FOR UP TO A 3X
LONGER LASTING
CLEAN FEELING*

***vs. brushing alone**

COOL MINT®

500 mL (1.05 Pt)

TARTAR CONTROL
KILLS GERMS

FIGHTS TARTAR BUILDUP & KILLS 99.9% OF GERMS

30039490

ANTISEPTIC

LISTERINE®

ULTRACLEAN

FOR UP TO A 3X LONGER LASTING CLEAN FEELING*

* vs. brushing alone

30039492

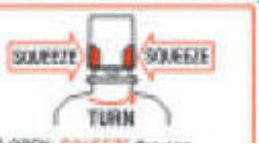
500 mL (1.05 Pt)

COOL MINT®

Drug Facts

Active ingredients — **Purposes**

Eucalyptol (0.092%)



LISTERINE ULTRACLEAN ANTISEPTIC COOL MINT

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

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0539
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 GREEN NO. 3 (UNII: 3P3ONR6O1S)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0539-2	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2012	
2	NDC:69968-0539-5	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2012	
3	NDC:69968-0539-1	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2012	
4	NDC:69968-0539-3	1500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part356	06/01/2012	

Labeler - Johnson & Johnson Consumer Inc. (002347102)

Revised: 5/2020 Johnson & Johnson Consumer Inc.