LISTERINE CLINICAL SOLUTIONS TEETH STRENGTH ALPINE MINT- sodium fluoride mouthwash
Johnson & Johnson Consumer Inc.

Listerine Clinical Solutions Teeth Strength Alpine Mint

Drug Facts

Active ingredient

Sodium Fluoride 0.02% (0.01% w/v Fluoride Ion)

Purpose

Anticavity

Uses

aids in the prevention of dental cavities

Warnings

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

Adults and children 12 years of age and older:

- use twice daily after brushing your teeth with a toothpaste
- vigorously swish 10 mL (2 teaspoonfuls) of rinse between your teeth for 1 minute and then spit out
- do not swallow the rinse
- do not eat or drink for 30 minutes after rinsing
- supervise children as necessary until capable of using without supervision

Children under 12 years of age: consult a dentist or a doctor

Other information

- store at room temperature
- cold weather may cloud this product

Inactive ingredients

Water, Alcohol, Sorbitol, Poloxamer 407, Flavor, Sodium Lauryl Sulfate, Eucalyptol, Sodium Saccharin, Methyl Salicylate, Thymol, Phosphoric Acid, Menthol, Sucralose, Disodium Phosphate, Green 3, Yellow 10

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

ANTICAVITY FLUORIDE MOUTHWASH

LISTERINE®

CLINICAL

SOLUTIONS

IMPORTANT: READ DIRECTIONS FOR PROPER USE

SODIUM FLUORIDE & ACIDULATED PHOSPHATE TOPICAL SOLUTION

TEETH STRENGTH

Repairs enamel now to prevent cavities

for 3X stronger* and longer-lasting teeth

(*with brushing in a lab study)

ALPINE MINT

1L (1 Qt 1.8 Fl Oz)

LISTERINE CLINICAL SOLUTIONS TEETH STRENGTH ALPINE MINT

sodium fluoride mouthwash

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0819
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	SODIUM FLUORIDE	0.1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
SORBITOL (UNII: 506T60A25R)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
WATER (UNII: 059QF0KO0R)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
EUCALYPTOL (UNII: RV6J6604TK)	
MENTHOL (UNII: L7T10EIP3A)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
THYMOL (UNII: 3J50XA376E)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0819-9	95 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/16/2023	
2	NDC:69968-0819-5	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/16/2023	
3	NDC:69968-0819-1	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	M021	10/16/2023	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 2/2024

Johnson & Johnson Consumer Inc.