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

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# Listerine Clinical Solutions Enamel 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 (21.6% v/v), 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

#### **ENAMEL STRENGTH**

Repairs enamel and prevents cavities

for stronger\*, longer lasting teeth

(\*with brushing in a lab study)

#### **ALPINE MINT**

1L (1 Qt 1.8 Fl Oz)



#### **3X STRONGER TEETH\***

30055979







## LISTERINE CLINICAL SOLUTIONS ENAMEL STRENGTH ALPINE MINT

sodium fluoride mouthwash

#### **Product Information** HUMAN OTC DRUG **Product Type Item Code (Source)** NDC:69968-0881 **Route of Administration ORAL**

#### **Active Ingredient/Active Moiety Ingredient Name Basis of Strength** Strength

SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU4080) SODIUM FLUORIDE 0.1 mg in 1 mL

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
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)	
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)	

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:69968- 0881-1	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/26/2024			
2	NDC:69968- 0881-9	95 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/26/2024			
3	NDC:69968- 0881-5	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/26/2024			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M021	02/26/2024		

# Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 3/2024 Johnson & Johnson Consumer Inc.