

**LISTERINE TOTAL CARE MILD ALCOHOL FREE ANTICAVITY FLUORIDE
MOUTHWASH- sodium fluoride mouthwash
Kenvue Brands LLC**

Listerine Total Care Mild Alcohol Free Anticavity Fluoride Mouthwash
Drug Facts

Active ingredient

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

Purpose

Anticavity

Use

aids in the prevention of dental cavities

Warnings

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 a day 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
- instruct children under 12 years of age in good rinsing habits (to minimize swallowing)
- supervise children as necessary until capable of using without supervision
- Children under 6 years of age: consult a dentist or a doctor

Other information

- Store at room temperature.

Inactive ingredients

Water, Sorbitol, Propylene Glycol, Sodium Lauryl Sulfate, Poloxamer 407, Flavor, Sodium Benzoate, Phosphoric Acid, Eucalyptol, Methyl Salicylate, Thymol, Sodium Saccharin, Menthol, Disodium Phosphate, Sucralose, Red 33, Blue 1

Questions?

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

Distributed by:

Kenvue Brands LLC

Summit, NJ 07901

PRINCIPAL DISPLAY PANEL - 1 L Bottle Label

ANTICAVITY FLUORIDE MOUTHWASH

LISTERINE®

TOTAL CARE

POWERED BY ESSENTIAL OILS + FLUORIDE

6

IN ONE

FOR A HEALTHIER MOUTH

360

CLEAN*

FRESHENS

BREATH

KILLS 99.9%

OF GERMS*

REPAIRS

ENAMEL

24 HR CAVITY

PROTECTION †

STRENGTHENS

TEETH

IMPORTANT: READ DIRECTIONS FOR PROPER USE

SODIUM FLUORIDE & ACIDULATED PHOSPHATE TOPICAL SOLUTION

1.0 L (1 Qt 1.8 FL Oz)



LISTERINE TOTAL CARE MILD ALCOHOL FREE ANTICAVITY FLUORIDE MOUTHWASH

sodium fluoride mouthwash

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0951
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
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
WATER (UNII: 059QF0KO0R)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
EUCALYPTOL (UNII: RV6J6604TK)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
SORBITOL (UNII: 506T60A25R)	
RED 33 (UNII: 9DBA0SBB0L)	

MENTHOL (UNII: L7T10EIP3A)				
DISODIUM PHOSPHATE (UNII: 22ADO53M6F)				
BLUE 1 (UNII: H3R47K3TBD)				
THYMOL (UNII: 3J50XA376E)				
SUCRALOSE (UNII: 96K6UQ3ZD4)				
SODIUM SACCHARIN (UNII: SB8ZUX40TY)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0951-1	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2026	
2	NDC:69968-0951-9	95 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2026	
3	NDC:69968-0951-5	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2026	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M021	02/01/2026	

Labeler - Kenvue Brands LLC (118772437)

Revised: 2/2026

Kenvue Brands LLC