LISTERINE TOTAL CARE FRESH MINT ANTICAVITY FLUORIDE- sodium fluoride mouthwash
Johnson & Johnson Consumer Inc.

Listerine Total Care Fresh Mint 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

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 using without supervision
- Children under 12 years of age: consult a dentist or doctor

Other information

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

Inactive Ingredients

Water, Alcohol (21.6% v/v),Sorbitol, Poloxamer 407,Eucalyptol,Flavor,Methyl Salicylate,Menthol,Phosphoric Acid,Sodium Saccharin

Thymol,Disodium Phosphate, Sucralose, Red 40, 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 - 1000 mL Bottle Label

ANTICAVITY FLUORIDE MOUTHWASH

LISTERINE®

TOTAL CARE

IMPORTANT: READ DIRECTIONS FOR PROPER USE.

SODIUM FLUORIDE & ACIDULATED PHOSPHATE TOPICAL SOLUTION

6 IN 1

BENEFITS

1. 24- HOUR CAVITY PROTECTION+
2. RESTORES ENAMEL
3. STRENGTHENS TEETH
4. FRESHENS BREATH
5. CLEANS THE WHOLE MOUTH
6. KILLS BAD BREATH GERMS

+When used twice daily * bad breath germs

FRESH MINT

1.0 L (1 Qt 1.8 fl oz)

LISTERINE TOTAL CARE FRESH MINT ANTICAVITY FLUORIDE

sodium fluoride mouthwash

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0787
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**
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
ALCOHOL (UNII: 3K9958V90M)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
EUCALYPTOL (UNII: RV6J6604TK)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	

THYMOL (UNII: 3J50XA376E)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0787-1	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/13/2023	
2	NDC:69968-0787-2	95 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/13/2023	
3	NDC:69968-0787-5	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/13/2023	
4	NDC:69968-0787-3	3 in 1 PACKAGE	01/13/2023	
4		1000 mL in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:69968-0787-4	2 in 1 PACKAGE	01/13/2023	
5		1000 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	01/13/2023	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 11/2023 Johnson & Johnson Consumer Inc.