LISTERINE SMART RINSE ANTICAVITY FLUORIDE RINSE - PINK LEMONADE FLAVOR- sodium fluoride mouthwash
Johnson & Johnson Consumer Inc.

LISTERINE ® SMART RINSE ® ANTICAVITY FLUORIDE RINSE PINK LEMONADE FLAVOR

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 6 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 doctor

Other information

- Store at room temperature
- Cold weather may temporarily cloud this product

Inactive ingredients

Water, Sorbitol, Flavor, Phosphoric Acid, Cetylpyridinium Chloride, Sucralose, Sodium

Saccharin, Disodium Phosphate, Red 40, Red 33

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

2X STRONGER TEETH FOR

BETTER CAVITY PROTECTION*

***with a brushing, in a lab study**

ALCOHOL FREE

ANTICAVITY FLUORIDE RINSE

LISTERINE ®

SMART RiNSE ®

SODIUM FLUORIDE & ACIDULATED PHOSPAHTE TOPICAL SOLUTION

PINK

LEMONADE

FLAVOR

500 mL (16.9Fl Oz)

IMPORTANT: Read

directions for proper use.

LISTERINE SMART RINSE ANTICAVITY FLUORIDE RINSE - PINK LEMONADE FLAVOR

sodium fluoride mouthwash

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0691
Route of Administration	ORAL		

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

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
CETYLPYRIDINIUM CHLORIDE (UNII: D9OM4SK49P)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0691-5	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	01/01/2021	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 4/2024

Johnson & Johnson Consumer Inc.