LISTERINE ESSENTIAL CARE- sodium monofluorophosphate gel, dentifrice
Johnson & Johnson Consumer Inc.

LISTERINE ® ESSENTIAL CARE ® GEL

Drug Facts

Active ingredient

Sodium monofluorophosphate 0.76% (0.13% w/v fluoride ion)

Purpose

Anticavity

Use

aids in the prevention of dental cavities

Warnings

Keep out of reach of children under 6 years of age. If more than used for brushing is accidently swallowed, get medical help or contact a Poison Control Center right away.

Directions

- adults and children 2 years of age and older: brush teeth thoroughly, preferably after each meal or at least 2 times a day, or as directed by a dentist or physician.
  - children 2-6 years use a pea sized amount
  - instruct children under 6 years of age in good brushing and rinsing habits (to minimize swallowing)
  - supervise children as necessary until capable of using without supervision.
- children under 2 years of age: ask a dentist or doctor

Other information

- Store at Room Temperature.

Inactive ingredients

water, sorbitol, hydrated silica, glycerin, PEG-32, sodium lauryl sulfate, cellulose gum, sodium saccharin, eucalyptol, methyl salicylate, thymol, phosphoric acid, menthol, sodium phosphate, xanthan gum, benzoic acid, flavor, mentha viridis (spearmint) leaf oil, disodium phosphate, blue 1, 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 - 119 g Tube Carton

LISTERINE ®

ESSENTIAL CARE ®

FLUORIDE ANTICAVITY TOOTHPASTE

ORIGINAL GEL
- **FRESHENS BREATH**
- **THE LISTERINE ® WAY**

POWERFULL MINT GEL

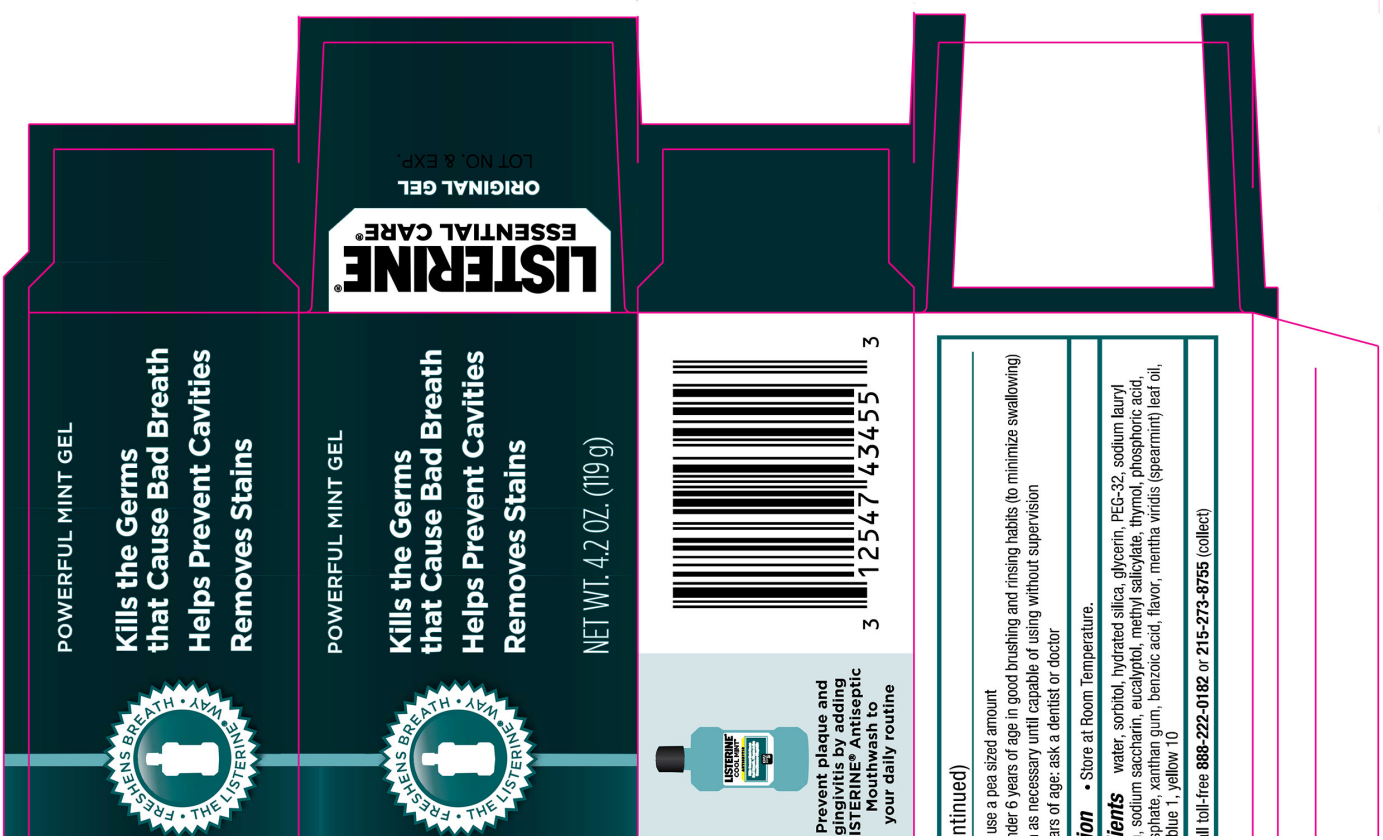
Kills the Germs

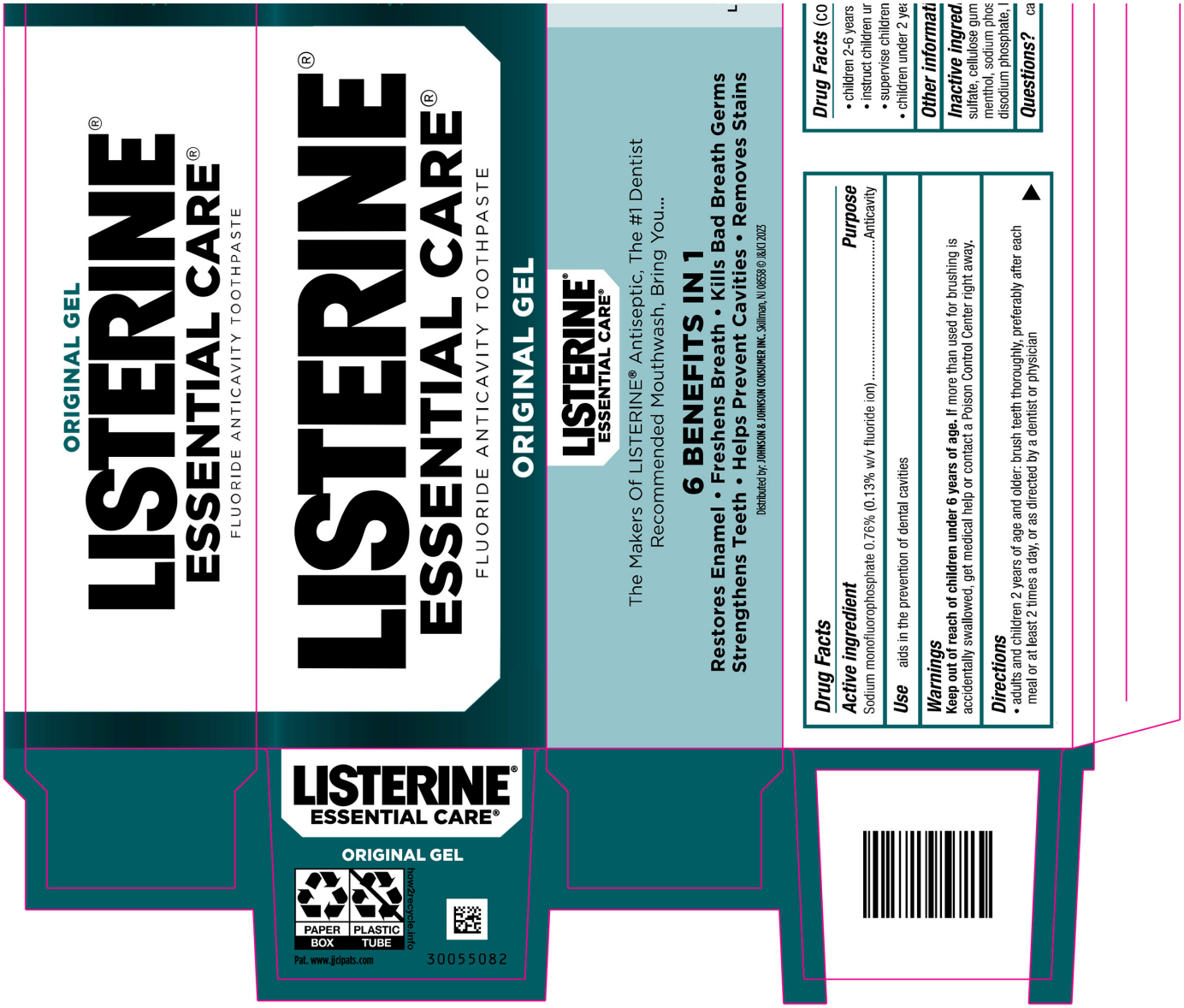
that Cause Bad Breath

Helps Prevent Cavities

Removes Stains

NET WT 4.2 OZ. (119 g)





LISTERINE ESSENTIAL CARE

sodium monofluorophosphate gel, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0701
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	1.3 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**

WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL 1600 (UNII: 1212Z7S33A)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
EUCALYPTOL (UNII: RV6J6604TK)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
THYMOL (UNII: 3J50XA376E)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
SODIUM PHOSPHATE (UNII: SE337SVY37)	
XANTHAN GUM (UNII: TTV12P4NEE)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
SPEARMINT OIL (UNII: C3M81465G5)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0701-4	1 in 1 CARTON	03/03/2021	
1		119 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M022	03/03/2021	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 10/2023 Johnson & Johnson Consumer Inc.