LISTERINE ORIGINAL ANTISEPTIC- eucalyptol, menthol, unspecified form, methyl salicylate, and thymol mouthwash
Kenvue Brands LLC

LISTERINE ® ORIGINAL ANTISEPTIC

Drug Facts

Active ingredient	***Purpose***
Eucalyptol (0.092%)	Antiplaque/antigingivitis
Menthol (0.042%)	Antiplaque/antigingivitis
Methyl Salicylate (0.060%)	Antiplaque/antigingivitis
Thymol (0.064%)	Antiplaque/antigingivitis

Use

helps prevent and reduce:
- plaque
- gingivitis

Warnings

Do not use in children under 12 years of age

Ask a dentist if symptoms persist, new symptoms appear, or conditions worsen after regular use

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
- rinse full strength for 30 seconds with 20mL (2/3 fluid ounce or 4 teaspoonfuls) morning and night
- do not swallow

Other information
- this rinse is not intended to replace brushing or flossing
- store at room temperature
- cold weather may cloud this product. Its antiseptic properties are not affected.

Inactive ingredients

water, alcohol (26.9% v/v), benzoic acid, poloxamer 407, sodium benzoate, caramel

Questions?

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

Distributed by:
Kenvue Brands LLC
Summit, NJ 07901

PRINCIPAL DISPLAY PANEL - 1.0 L Bottle Label

ANTIGINGIVITIS / ANTIPLAQUE MOUTHWASH

LISTERINE®

ORIGINAL

ANTISEPTIC

FOR A FRESHER &

CLEANER MOUTH THAN

BRUSHING ALONE

1.0 L (1 Qt 1.8 Fl Oz)

LISTERINE ORIGINAL ANTISEPTIC

eucalyptol, menthol, unspecified form, methyl salicylate, and thymol mouthwash

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0400
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
EUCALYPTOL (UNII: RV6J6604TK) (EUCALYPTOL - UNII:RV6J6604TK)	EUCALYPTOL	0.92 mg in 1 mL
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	0.42 mg in 1 mL
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	0.6 mg in 1 mL
THYMOL (UNII: 3J50XA376E) (THYMOL - UNII:3J50XA376E)	THYMOL	0.64 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
CARAMEL (UNII: T9D99G2B1R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0400-2	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/06/2019	09/02/2023
2	NDC:69968-0400-5	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/06/2019	10/01/2023
3	NDC:69968-0400-3	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/06/2019	
4	NDC:69968-0400-1	1500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/06/2019	
5	NDC:69968-0400-9	95 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/06/2019	10/07/2021
6	NDC:69968-0400-8	2 in 1 PACKAGE	01/06/2019	
6	NDC:69968-0400-1	1500 mL in 1 BOTTLE, PLASTIC; 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	01/06/2019	

Labeler - Kenvue Brands LLC (118772437)