SUPER VANILLA FLUORIDE MOUTHWASH- anticavity mouthwash liquid
SuperMouth, LLC.

Super Vanilla Fluoride Mouthwash

ACTIVE INGREDIENT
(in each 10 mL)

Sodium Fluoride 0.02%

PURPOSE
Sodium Fluoride …. Anticavity

USE

Fluoride mouthwash

WARNINGS:

Keep out of reach of children under 6 years of age.

If more than used for rinsing is accidentally swallowed, get medical help or contact a Poison Control Center right away.

Do not use if safety seal is broken.

DIRECTIONS

1) Shake well before using.
2) Use daily by dispensing 10 mL, swish vigorously for **1 minute**, and then spit out.
3) No need to rinse.

INACTIVE INGREDIENTS

Water, Erythritol, Xylitol, *Hydroxamin® (Nano-Hydroxyapatite, Menaquinone-7 (Vit K2), Cholecalciferol (Vit D3)), Inulin, Glycerin, Sodium Gluconate, Natural Flavors, Sodium Bicarbonate, Methyl -sulfonylmethane, Natural Benzoic Acid, Sodium Ascorbate (Vit C), Xanthan Gum.

Product Labeling: Super Vanilla Fluoride Mouthwash

Supermouth

super vanilla

fluoride mouthwash

reaches areas brushing & flossing don't and prevents bad breath

16 fl oz (473 mL)

SuperMouth®
9737 Aero Drive
San Diego, CA 92123

(844) MOUTHCARE

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SUPER VANILLA FLUORIDE MOUTHWASH

anticavity mouthwash liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83729-104
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	0.002 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ERYTHRITOL (UNII: RA96B954X6)	
XYLITOL (UNII: VCQ006KQ1E)	
TRIBASIC CALCIUM PHOSPHATE (UNII: 91D9GV0Z28)	
MENAQUINONE 7 (UNII: 8427BML8NY)	
CHOLECALCIFEROL (UNII: 1C6V77QF41)	
INULIN (UNII: JOS53KRJ01)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM GLUCONATE (UNII: R6Q3791S76)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
SODIUM ASCORBATE (UNII: S033EH8359)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	gray	**Score**	
Shape		**Size**	
Flavor	VANILLA	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83729-104-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/21/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	03/21/2023	

Labeler - SuperMouth, LLC. (049384038)

Registrant - SuperMouth, LLC. (049384038)