SUPER MINT PRESCRIPTION- anticavity toothpaste paste, dentifrice
SuperMouth, LLC.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

SuperMouth Super Mint Prescription Toothpaste

ACTIVE INGREDIENT
Sodium Fluoride 1.1% (5000 ppm)

PURPOSE
Sodium Fluoride .... Anticavity

USE: This medication is used as part of a professional program for the prevention and control of dental caries. Use as directed by your health professional.

WARNINGS: Do not swallow. Do not use in children under 6 years of age unless recommended by a dentist or physician. Keep out of reach of children under 6 years of age. If more than used for brushing is accidentally swallowed, get medical help or contact a Poison Control Center right away.

See your dentist if the problem persists or worsens. Do not use this product longer than 4 weeks unless recommended by a dentist or physician.

DIRECTIONS: Use as directed by a dentist or physician.

INACTIVE INGREDIENTS

Glycerin, Water, Hydrated Silica, *Hydroxamin® (Nano-Hydroxyapatite, Menquinone-y (Vit K2), Cholecalciferol (Vit D3)), Xylitol, Inulin, Sodium Gluconate, Methylsulfonylmethane, Xanthan Gum, Quillaja Saponaria Extract, Natural Flavor, Natural Benzoic Acid, Sodium Ascorbate (Vit C), Stevia Leaf Extract, Cranberry Seed Oil.
*Patent-Pending Formulation

PRINCIPAL DISPLAY PANEL - SuperMouth Super Mint Prescription Toothpaste

Supermouth Pro

1.1% sodium fluoride

5000 ppm

Anticavity Toothpaste

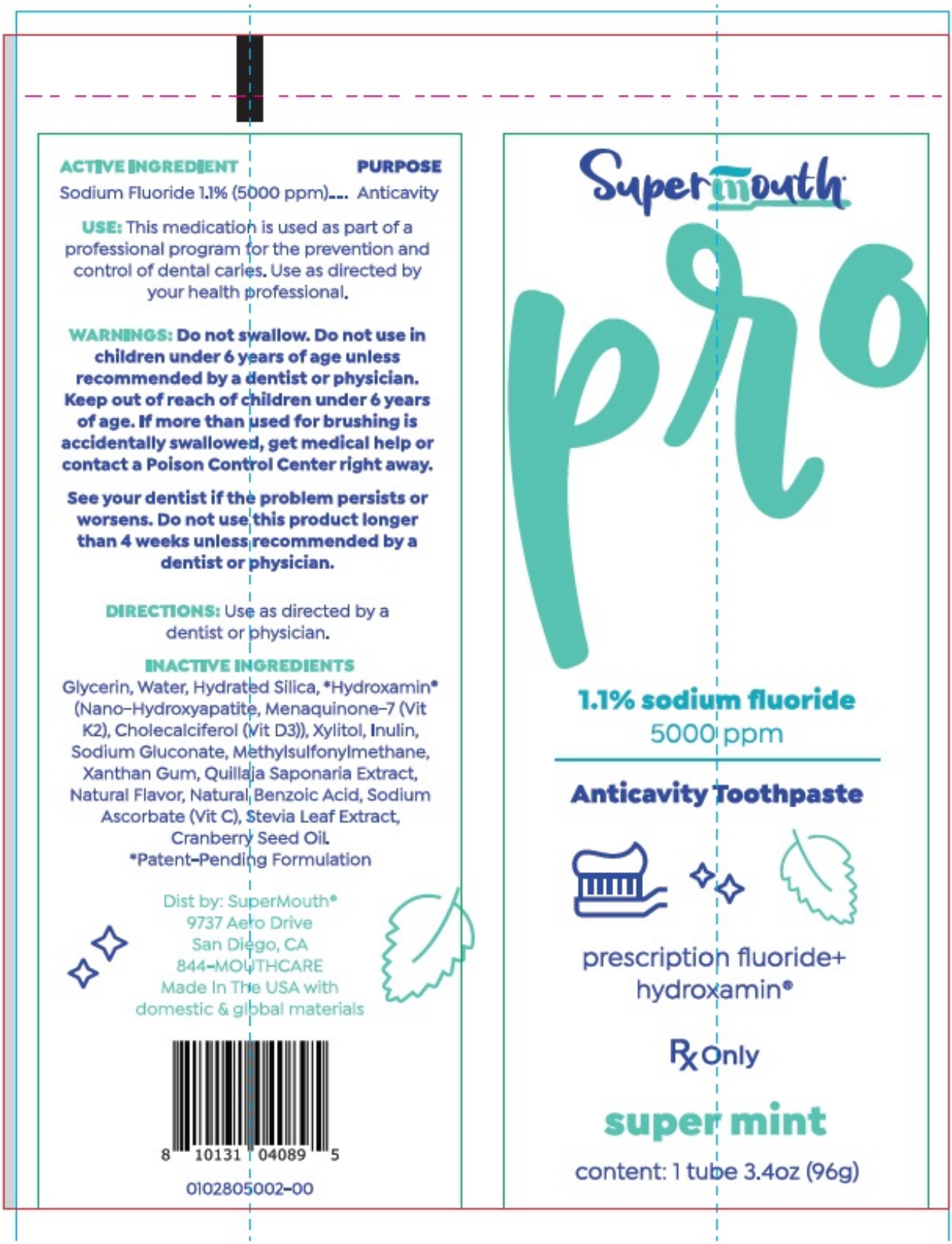
Rx Only

super mint

content: 1 tube 3.4oz (96g)

Dist by: SuperMouth®
9737 Aero Drive
San Diego, CA

844-MOUTHCARE
Made In The USA with
Domestic & global materials



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SUPER MINT PRESCRIPTION
anticavity toothpaste paste, dentifrice

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:83729-102
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	1.1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
TRIBASIC CALCIUM PHOSPHATE (UNII: 91D9GV0Z28)	
MENAQUINONE 7 (UNII: 8427BML8NY)	
CHOLECALCIFEROL (UNII: 1C6V77QF41)	
XYLITOL (UNII: VCQ006KQ1E)	
INULIN (UNII: JOS53KRJ01)	
SODIUM GLUCONATE (UNII: R6Q3791S76)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
XANTHAN GUM (UNII: TTV12P4NEE)	
QUILLAJA SAPONARIA WHOLE (UNII: HIU9R169Y7)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
SODIUM ASCORBATE (UNII: S033EH8359)	
STEVIA LEAF (UNII: 6TC6NN0876)	
CRANBERRY SEED OIL (UNII: 73KDS3BW5E)	

Product Characteristics

Color	brown	**Score**	
Shape		**Size**	
Flavor	MINT	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83729-102-10	1 in 1 CARTON	10/13/2023	
1		96 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
unapproved drug other		10/13/2023	

Labeler - SuperMouth, LLC. (049384038)

Registrant - SuperMouth, LLC. (049384038)

Revised: 11/2025 SuperMouth, LLC.