# 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.

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#### 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

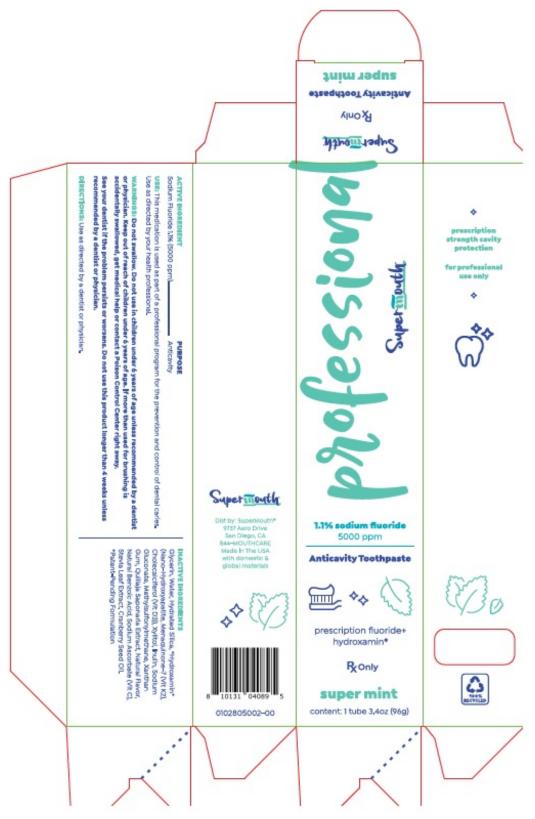
Rx Only

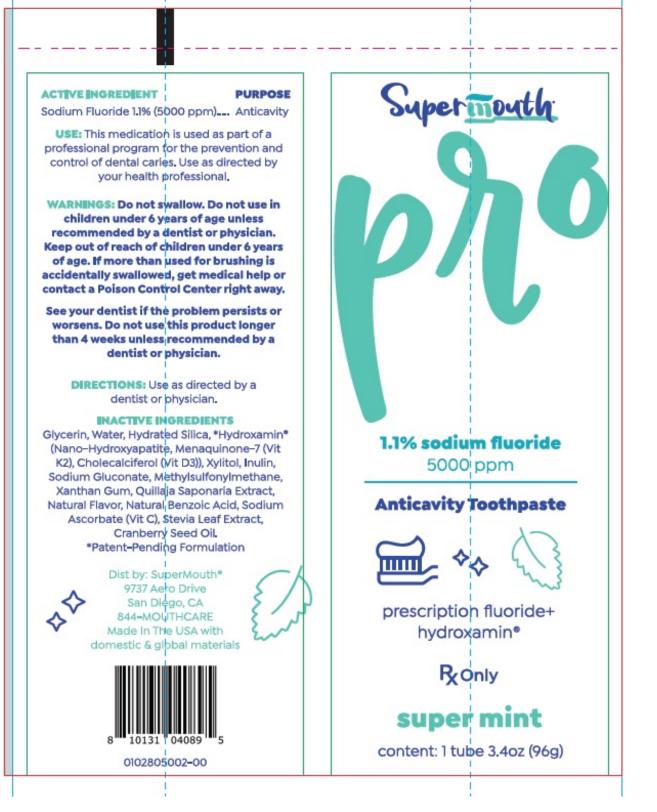
super mint

#### content: 1 tube 3.4oz (96g)

Dist by: SuperMouth<sup>®</sup> 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 Inform				H				
Product Type		HUMAN PRESCRIPTIC	IN DRUG	Item Code (Source) NDC		2:83729-102		
Route of Adminis	stration	DENTAL						
Active Ingredie	ent/Active	Moiety						
Ingredient Name Basis of Stru						enath	Strength	
	474W7) (FLUORIDE ION	I - UNII:Q80V	PU408O)	FLUORIDE ION	_	1.1 g in 100		
							J	
nactive Ingree	dients							
		Ingredient Nar	ne			9	Strength	
GLYCERIN (UNII: PD	C6A3C0OX)							
VATER (UNII: 059QF	OKOOR)							
HYDRATED SILICA	(UNII: Y607T40	G8P9)						
RIBASIC CALCIUM	I PHOSPHATE	(UNII: 91D9GV0Z28)						
MENAQUINONE 7 (	UNII: 8427BML	8NY)						
CHOLECALCIFERO	L (UNII: 1C6V7	7QF41)						
XYLITOL (UNII: VCQ	006KQ1E)							
NULIN (UNII: JOS53	KRJ01)							
SODIUM GLUCONA	TE (UNII: R6Q3	3791576)						
DIMETHYL SULFON	<b>IE</b> (UNII: 9H4PC	04Z4FT)						
KANTHAN GUM (UN	II: TTV12P4NEE	E)						
QUILLAJA SAPONAF	RIA WHOLE (U	NII: HIU9R169Y7)						
BENZOIC ACID (UNI	II: 85KN0B0MIN	1)						
SODIUM ASCORBA	<b>TE</b> (UNII: S033	EH8359)						
STEVIA LEAF (UNII:	6TC6NN0876)							
CRANBERRY SEED	OIL (UNII: 73K	DS3BW5E)						
Product Chara	cteristics							
Color		brown	Score					
Shape			Size					
Flavor		MINT	Imprint Code					
Contains			imprint c	oue				
contains								
Packaging								
# Item Code	Pac	ckage Descriptio	n		ting Start Date		eting End Date	
<b>1</b> NDC:83729-102-	1 in 1 CARTON	J		10/13/2023			Dute	
<b>1</b> 0 <b>1</b>	96 g in 1 TUBE; Type 0: Not a Combination							
	Product							
Marketing I	nformat	ion						

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

# Labeler - SuperMouth, LLC. (049384038)

## Registrant - SuperMouth, LLC. (049384038)

Revised: 10/2023

SuperMouth, LLC.