MADE BY DENTISTS ENAMEL REBUILDING FRESH MINT- potassium nitrate, sodium fluoride paste MADE BY DENTISTS INC

Made by Dentists Enamel Rebuilding Toothpaste Fresh Mint

Drug Facts

Active ingredient:

Potassium Nitrate 5.0% Sodium Fluoride 0.24% (0.15% w/v fluoride ion)

Purpose

Antisensitivity

Anticavity

Use:

Aids in the prevention of cavities. Helps reduce painful sensitivity of the teeth to cold, heat, acids, sweets, or contact.

Warnings:

Sensitive teeth may indicate a serious problem that may need prompt care by a dentist. See your dentist if the problem persists or worsens. Do not use this product longer than 4 weeks unless recommended by a dentist or doctor.

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 12 years of age and older: Apply at least a 1-inch strip of the product onto a soft bristle toothbrush. Brush teeth thoroughly, for at least 1 minute twice a day, (morning and evening) or as recommended by a dentist or doctor.
- Make sure to brush all sensitive areas of the teeth.
- Children under 12 years of age: Consult a dentist or doctor.

Inactive ingredients:

Cellulose Gum, Cocamidopropyl Betaine, Flavor, Hydrated Silica, Hydroxyapatite, Pentasodium Triphosphate, Potassium Sorbate, Sodium Bicarbonate, Sodium Saccharin,

Questions or comments?

+ 1 (646) 980-6461

Label







Made By Dentists[™]

ENAMEL REBUILDING PROFESSIONAL TOOTHPASTE

Fluoride Anticavity Toothpaste for Sensitive Teeth NET WT 4.2 Oz / 120 g

Professional Oral Care

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Sodium Fluoride 0.24% Anticavity
(0.15% w/v fluoride ion)

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- × Gluten Free × Paraben Free
- × Sulfate Free× Phthalate Free× Cruelty Free× Vegan Friendly





FRESH MINT FLAVOR

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MADE BY DENTISTS ENAMEL REBUILDING FRESH MINT

potassium nitrate, sodium fluoride paste

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:75065-028

Route of Administration DENTAL

telli code (30di)

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
POTASSIUM NITRATE (UNII: RU45X2JN0Z) (NITRATE ION - UNII:T93E9Y2844)	POTASSIUM NITRATE	50 mg in 1 g		
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU4080)	FLUORIDE ION	1.5 mg in 1 g		

Inactive Ingredients		
Ingredient Name	Strength	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311)		
COCAMIDOPROPYL BETAINE (UNII: 50CF3011KX)		
HYDRATED SILICA (UNII: Y6O7T4G8P9)		
TRIBASIC CALCIUM PHOSPHATE (UNII: 91D9GV0Z28)		
SODIUM TRIPOLYPHOSPHATE ANHYDROUS (UNII: 9SW4PFD2FZ)		
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)		
SODIUM BICARBONATE (UNII: 8MDF5V39QO)		
SACCHARIN SODIUM (UNII: SB8ZUX40TY)		
SORBITOL (UNII: 506T60A25R)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		
WATER (UNII: 059QF0KO0R)		
XYLITOL (UNII: VCQ006KQ1E)		

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:75065-028- 06	1 in 1 CARTON	02/22/2024				
1		120 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	02/22/2024			

Labeler - MADE BY DENTISTS INC (117405870)

Revised: 2/2024 MADE BY DENTISTS INC