

ARC WHITENING- sodium fluoride paste, dentifrice
Procter & Gamble Manufacturing Company

ARC WHITENING FLUORIDE TOOTHPASTE CHARCOAL

Drug Facts

Active ingredient

Sodium fluoride 0.243% (0.14% w/v fluoride ion)

Purpose

Anticavity toothpaste

Use

helps protect against cavities

Warnings

Keep out of reach of children under 6 yrs of age. If more than used for brushing is accidentally swallowed, get medical help or contact a Poison Control Center right away.

Directions

- adults and children 2 yrs. & older: brush teeth thoroughly after meals or at least twice a day or use as directed by a dentist
- do not swallow
- to minimize swallowing use a pea-sized amount in children under 6
- supervise children's brushing until good habits are established
- children under 2 yrs.: ask a dentist

Inactive ingredients

glycerin, water, hydrated silica, xylitol, flavor, mentha piperita (peppermint) oil, charcoal powder, sodium cocoyl glutamate, xanthan gum, carrageenan, cocamidopropyl betaine, stevia rebaudiana extract

Questions?

1-866-989-3951

DISTR. BY PROCTER & GAMBLE, CINCINNATI, OH 45202

ARC WHITENING

sodium fluoride paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69423-889
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	1.4 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ACTIVATED CHARCOAL (UNII: 2P3VWUJ3H10)	
CARRAGEENAN (UNII: 5C69YCD2YJ)	
XANTHAN GUM (UNII: TTV12P4NEE)	
WATER (UNII: 059QF0KO0R)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
GLYCERIN (UNII: PDC6A3C0OX)	
STEVIA REBAUDIANA WHOLE (UNII: 6U422Y08O2)	
SODIUM COCOYL GLUTAMATE (UNII: BMT4RCZ3HG)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
XYLITOL (UNII: VCQ006KQ1E)	

Product Characteristics

Color	black	Score	
Shape		Size	
Flavor	MINT	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-889-40	1 in 1 CARTON	01/05/2021	
1		113 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
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OTC Monograph Drug M021

01/05/2021

Labeler - Procter & Gamble Manufacturing Company (004238200)

Revised: 10/2023

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