

CINNAFRESH- sodium fluoride gel, dentifrice
Bob Barker Company Inc.

CinnaFresh

Drug Facts

ACTIVE INGREDIENT

Sodium Fluoride - 0.22% (0.1% W/V fluoride ion)

PURPOSE

Anticavity toothpaste

Use

Helps protect against cavities.

Warnings

Keep out of the 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 immediately.

Directions

Adults & Children 6 years of age & older:	Brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or doctor.
Children 2 to 6 years:	Use only a pea sized amount and supervise child's brushing and rinsing (to minimize swallowing).
Children under 2 years:	Ask a dentist or physician.

Inactive ingredients

Water, Sorbitol, Carbomer, Sodium Lauryl Sulfate, Flavor, Sodium Hydroxide, Sodium Saccharin, Disodium EDTA, Sodium Benzoate, Red 40

Dist. by Bob Barker Co. Inc. Fuquay-Varina, NC 27526

PRINCIPAL DISPLAY PANEL - 24 GRAMS Tube Label

CinnaFresh
 anticavity gel toothpaste
 Sodium Fluoride 0.22%
 Bold Cinnamon Flavor
 NET WT. 0.85 OZ
 (24 GRAMS)

CinnaFresh

anticavity gel toothpaste

Sodium Fluoride 0.22%

Bold Cinnamon Flavor

**NET WT. 0.85 OZ
(24 GRAMS)**

Drug Facts:	Drug Facts (continued)	
ACTIVE INGREDIENT: PURPOSE: Sodium Fluoride - 0.22% (0.1% W/V fluoride ion)...Anticavity toothpaste	Directions: Adults & Children 6 years of age & older:	Brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or doctor.
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Batch no., mfg. date & exp. date on crimp.

E.C. No.

GUJ/COS/GC/1918



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Made in India Dist. by Bob Barker Co. Inc. Fuquay-Varina, NC 27526

CINNAFRESH			
sodium fluoride gel, dentifrice			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53247-133
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Sodium Fluoride (UNII: 8ZYQ1474W7) (Fluoride Ion - UNII:Q80VPU4080)	Sodium Fluoride	2.2 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Sorbitol (UNII: 506T60A25R)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
Sodium Lauryl Sulfate (UNII: 368GB5141J)	
Sodium Hydroxide (UNII: 55X04QC32I)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
Sodium Benzoate (UNII: OJ245FE5EU)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53247-133-01	24 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	M021	01/01/2024	

Labeler - Bob Barker Company Inc. (058525536)

Revised: 4/2024

Bob Barker Company Inc.