CREST PRO-HEALTH ADVANCED ANTI-CAVITY MAX CAVITY- sodium fluoride rinse
The Procter & Gamble Manufacturing Company

Crest ®
Pro-Health Advanced
Anti-Cavity
Max Cavity Protection

Drug Facts

Active ingredient

Sodium Fluoride 0.02% (0.01% w/v fluoride ion)

Purpose

Anticavity

Use

Aids in the prevention of dental cavities

Warnings

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

Directions

- Adults and children 6 years & older:
  Use twice a day after brushing your teeth with a toothpaste.
- Vigorously swish 10 mL (2 teaspoonfuls) of rinse between your teeth for 1 minute and then spit out.
- Do not swallow the rinse.
- Do not eat or drink for 30 minutes after rinsing.
- Instruct children under 12 years of age in good rinsing habits (to minimize swallowing).
- Supervise children as necessary until capable of using without supervision.
- Children under 6 years of age: Consult a dentist or doctor.

Inactive ingredients

water, glycerin, flavor, poloxamer 407, cetylpyridinium chloride, sodium saccharin, phosphoric acid, methylparaben, sucralose, propylparaben, disodium phosphate, yellow 6, green 3

Questions?

1–800–285–9139

DISTR. BY PROCTER & GAMBLE,

CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 500 mL Bottle Label

+ FLUORIDE

Crest

PRO-HEALTH

ADVANCED

ANTICAVITY FLUORIDE MOUTHWASH

ALCOHOL FREE

ANTI-CAVITY

IMPORTANT:

Read directions for proper use

MAX CAVITY

PROTECTION

PROTECTS AGAINST CAVITIES

REBUILDS WEAKENED ENAMEL

STRENGTHENS TEETH

CLEANS THE WHOLE MOUTH

FRESHENS BREATH

NO ALCOHOL BURN

500 mL

(16.9 fl oz)

CREST PRO-HEALTH ADVANCED ANTI-CAVITY MAX CAVITY

sodium fluoride rinse

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69423-848
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	2.2 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
CETYLPYRIDINIUM CHLORIDE (UNII: D9OM4SK49P)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	

PHOSPHORIC ACID (UNII: E4GA8884NN)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-848-50	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/31/2020	
2	NDC:69423-848-10	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/31/2020	
3	NDC:69423-848-01	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/31/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	01/31/2020	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 4/2024 The Procter & Gamble Manufacturing Company