CREST PRO-HEALTH ADVANCED W/EXTRA WHITENING- sodium fluoride rinse The Procter & Gamble Manufacturing Company

Crest [®] Pro-Health™

Advanced w/Extra Whitening

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 12 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 12 years of age: consult a dentist or doctor.

Inactive ingredients

water, glycerin, hydrogen peroxide, poloxamer 407, flavor, polysorbate 80, sodium saccharin, sucralose, phosphoric acid, disodium phosphate

Questions?

1-800-285-9139

DISTR. BY PROCTER & GAMBLE, CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 946 mL Bottle Label

ALCOHOL FREE WITH FLUORIDE

Crest[®] PRO-HEALTH[™] ADVANCED ANTICAVITY FLUORIDE MOUTHWASH

STRONGER TEETH* FOR A HEALTHIER MOUTH

WITH EXTRA WHITENING

- KILLS BAD BREATH GERMS
- WHITER SMILE** IN 7 DAYS
- HELPS PREVENT CAVITIES
- STRENGTHENS ENAMEL
- FRESHENS BREATH

ENERGIZING MINT

IMPORTANT: Read directions for proper use.

946 mL (32 FL OZ)

97275245



CREST PRO-HEALTH ADVANCED W/EXTRA WHITENING

sodium fluoride rinse

Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC		NDC:	C:37000-867	
Route of Administration	ORAL					
Active Ingredient/Active	e Moietv					
Ingredient Name Basis of Strength						
				Strength		
SODIUM FLUORIDE (UNII: 8ZYQ		II:Q80VPU408O)		5	0.1 mg in 1 mL	
SODIUM FLUORIDE (UNII: 8ZYQ		II:Q80VPU408O)		J	0.1 mg in 1 mL	
SODIUM FLUORIDE (UNII: 8ZYQ Inactive Ingredients		II:Q80VPU408O)		5	0.1 mg in 1 mL	
		II:Q80VPU408O)		J	0.1 mg in 1 mL Strength	
	1474W7) (FLUORIDE ION - UN	II:Q80VPU408O)			-	
Inactive Ingredients	1474W7) (FLUORIDE ION - UN	II:Q80VPU408O)			-	
Inactive Ingredients WATER (UNII: 059QF0KO0R)	1474W7) (FLUORIDE ION - UN Ingredient Name	II:Q80VPU408O)			-	
Inactive Ingredients WATER (UNII: 059QF0KO0R) GLYCERIN (UNII: PDC6A3C0OX)	1474W7) (FLUORIDE ION - UN Ingredient Name 8X060AN9V)	II:Q80VPU408O)			-	

SA	CCHARIN SOD	IUM (UNII: S	B8ZUX40TY)				
รบ	CRALOSE (UNI	I: 96K6UQ3Z	D4)				
P۲	IOSPHORIC AC	ID (UNII: E40	GA8884NN)				
sc	DIUM PHOSPH	IATE, DIBAS	SIC, ANHYDROUS (UNII: 22ADO53	M6F)			
P	roduct Chai	racteristi	cs				
Color			white (Clear)	Scor	Score		
Shape				Size	ze		
Flavor			MINT	Impr	orint Code		
Contains							
_							
Pa	ackaging						
#	Item Code		Package Description		Marketing Start Date	Marketing End Date	
1	NDC:37000- 867-01	473 mL in 1 Combinatio	. BOTTLE, PLASTIC; Type 0: Not a n Product		07/20/2015		
2	NDC:37000- 867-02	946 mL in 1 Combinatio	. BOTTLE, PLASTIC; Type 0: Not a n Product		07/20/2015		
Μ	arketing	Inform	nation				
	Marketing Category	Арр	lication Number or Monogra Citation	ph	Marketing Start Date	Marketing End Date	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 10/2023

The Procter & Gamble Manufacturing Company