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

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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:37000-867
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	0.1 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	

SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	

Product Characteristics

Color	white (Clear)	**Score**	
Shape		**Size**	
Flavor	MINT	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37000-867-01	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/2015	
2	NDC:37000-867-02	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part355	07/20/2015	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 5/2023 The Procter & Gamble Manufacturing Company